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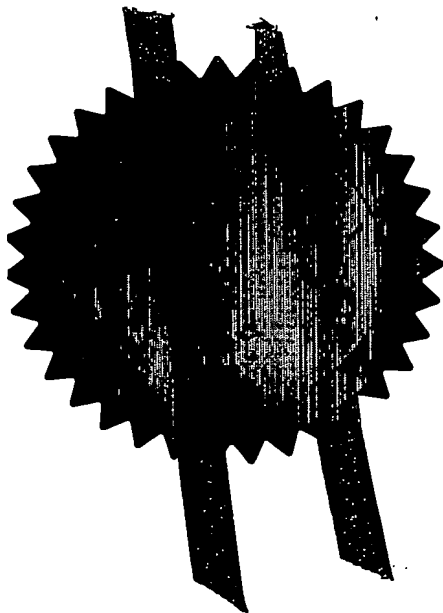
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Dated

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Request for grant of a patent

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The Patent Office

Cardiff Road
Newport
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1. Your reference

100660

2. Patent application number

(The Patent Office will fill in this part)

0205165.4

- 6 MAR 2002

3. Full name, address and postcode of the or of each applicant (underline all surnames)

AstraZeneca AB
S-151 85 Sodertalje
Sweden

Patents ADP number (if you know it)

1822448003

If the applicant is a corporate body, give the country/state of its incorporation

Sweden

4. Title of the invention

CHEMICAL COMPOUNDS

5. Name of your agent (if you have one)

Rachel M. Tinsley

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

AstraZeneca UK Limited
Global Intellectual Property
Mercedes, Alderley Park
Macclesfield
Cheshire SK10 4TG

Patents ADP number (if you know it)

6921795002

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)

Date of filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

- a) any applicant named in part 3 is not an inventor, or
 - b) there is an inventor who is not named as an applicant, or
 - c) any named applicant is a corporate body.
- See note (d)

Patents Form 1/77

9. Enter the number of sheets for any of the following items you are filing with this form. Do not count copies of the same document

Continuation sheets of this form

Description

Claim(s)

Abstract

Drawing(s)

64

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for preliminary examination and search (Patents Form 9/77)

Request for substantive examination (Patents Form 10/77)

Any other documents (please specify)

11.

I/We request the grant of a patent on the basis of this application.

Signature

Date

Authorised Signatory

05/03/2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Joanne M. Marshall - 01625 - 516485

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CHEMICAL COMPOUNDS

The present invention relates to heterocyclic amide derivatives, pharmaceutically acceptable salts and *in vivo* hydrolysable esters thereof. These heterocyclic amides possess glycogen phosphorylase inhibitory activity and accordingly have value in the treatment of disease states associated with increased glycogen phosphorylase activity and thus are potentially useful in methods of treatment of a warm-blooded animal such as man. The invention also relates to processes for the manufacture of said heterocyclic amide derivatives, to pharmaceutical compositions containing them and to their use in the manufacture of medicaments to inhibit glycogen phosphorylase activity in a warm-blooded animal such as man.

The liver is the major organ regulating glycaemia in the post-absorptive state. Additionally, although having a smaller role in the contribution to post-prandial blood glucose levels, the response of the liver to exogenous sources of plasma glucose is key to an ability to maintain euglycaemia. An increased hepatic glucose output (HGO) is considered to play an important role in maintaining the elevated fasting plasma glucose (FPG) levels seen in type 2 diabetics; particularly those with a FPG >140mg/dl (7.8mM). (Weyer et al, (1999), J Clin Invest 104: 787-794; Clore & Blackgard (1994), Diabetes 43: 256-262; De Fronzo, R. A., et al, (1992) Diabetes Care 15; 318 - 355; Reaven, G.M. (1995) Diabetologia 38; 3-13).

Since current oral, anti-diabetic therapies fail to bring FPG levels to within the normal, non-diabetic range and since raised FPG (and glycHbA1c) levels are risk factors for both macro- (Charles, M.A. et al (1996) Lancet 348, 1657-1658; Coutinho, M. et al (1999) Diabetes Care 22; 233-240; Shaw, J.E. et al (2000) Diabetes Care 23, 34-39) and micro-vascular disease (DCCT Research Group (1993) New. Eng. J. Med. 329; 977-986); the reduction and normalisation of elevated FPG levels remains a treatment goal in type 2 DM.

It has been estimated that, after an overnight fast, 74% of HGO was derived from glycogenolysis with the remainder derived from gluconeogenic precursors (Hellerstein et al (1997) Am J Physiol, 272: E163). Glycogen phosphorylase is a key enzyme in the generation by glycogenolysis of glucose-1-phosphate, and hence glucose in liver and also in other tissues such as muscle and neuronal tissue.

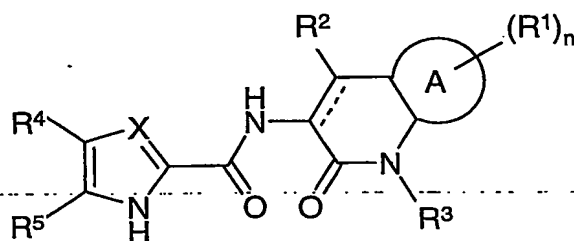
Liver glycogen phosphorylase a activity is elevated in diabetic animal models including the db/db mouse and the fa/fa rat (Aiston S et al (2000). Diabetologia 43, 589-597).

Inhibition of hepatic glycogen phosphorylase with chloroindole inhibitors (CP91149 and CP320626) has been shown to reduce both glucagon stimulated glycogenolysis and glucose output in hepatocytes (Hoover et al (1998) J Med Chem 41, 2934-8; Martin et al (1998) PNAS 95, 1776-81). Additionally, plasma glucose concentration is reduced, in a dose related manner, db/db and ob/ob mice following treatment with these compounds.

Studies in conscious dogs with glucagon challenge in the absence and presence of another glycogen phosphorylase inhibitor, Bay K 3401, also show the potential utility of such agents where there is elevated circulating levels of glucagon, as in both Type 1 and Type 2 diabetes. In the presence of Bay R 3401, hepatic glucose output and arterial plasma glucose following a glucagon challenge were reduced significantly (Shiota et al, (1997), Am J Physiol, 273: E868).

The heterocyclic amides of the present invention possess glycogen phosphorylase inhibitory activity and accordingly are expected to be of use in the treatment of type 2 diabetes, insulin resistance, syndrome X, hyperinsulinaemia, hyperglucagonaemia, cardiac ischaemia and obesity, particularly type 2 diabetes.

According to one aspect of the present invention there is provided a compound of formula (1):



(1)

wherein:

---- is a single or double bond;

X is N or CH;

R^4 and R^5 together are either $-S-C(R^6)=C(R^7)-$ or $-C(R^7)=C(R^6)-S-$;

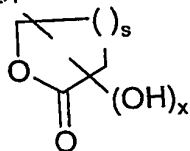
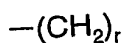
- wherein R^6 and R^7 are independently selected from hydrogen, halo, nitro, cyano, hydroxy, fluoromethyl, difluoromethyl, trifluoromethyl, trifluoromethoxy, amino, carboxy, carbamoyl, mercapto, sulfamoyl, ureido, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{1-6} alkoxy, C_{1-6} alkanoyl, C_{1-6} alkanoyloxy, $N-(C_{1-6}alkyl)amino$, $N,N-(C_{1-6}alkyl)_2amino$, $C_{1-6}alkanoylamino$, $N-(C_{1-6}alkyl)carbamoyl$, $N,N-(C_{1-6}alkyl)_2carbamoyl$, $C_{1-6}alkylS(O)_a$ wherein a is 0 to 2, $C_{1-6}alkoxycarbonyl$, $C_{1-6}alkoxycarbonylamino$, $N-(C_{1-6}alkyl)sulphamoyl$, $N,N-(C_{1-6}alkyl)_2sulphamoyl$, $C_{1-6}alkylsulphonylamino$ and $C_{1-6}alkylsulphonyl-N-(C_{1-6}alkyl)amino$;
- wherein:

A is phenylene or heteroarylene;

- 15 n is 0, 1 or 2;

wherein R^1 is independently selected from hydrogen, halo, nitro, cyano, hydroxy, amino, carboxy, carbamoyl, $N-C_{1-4}alkylcarbamoyl$, $N,N-(C_{1-4}alkyl)_2carbamoyl$, sulphamoyl, $N-C_{1-4}alkylsulphamoyl$, $N,N-(C_{1-4}alkyl)_2sulphamoyl$, sulfinio, sulfo, $C_{1-4}alkyl$, $C_{2-4}alkenyl$, $C_{2-4}alkynyl$, $C_{1-4}alkoxy$, $C_{1-4}alkanoyl$, $C_{1-4}alkanoyloxy$, $N-(C_{1-4}alkyl)amino$, $N,N-(C_{1-4}alkyl)_2amino$, hydroxy $C_{1-4}alkyl$, fluoromethyl, difluoromethyl, trifluoromethyl, trifluoromethoxy, $C_{1-4}alkoxy$ and

20 R^1 is of the formula A' or A'':



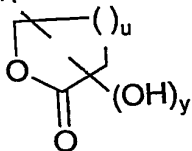
(A')



- 25 wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

R^2 is hydrogen, hydroxy or carboxy;

R^3 is selected from hydrogen, hydroxy, C_{1-4} alkanoyl, carbamoyl, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), C_{5-7} cycloalkyl (optionally substituted with 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano(C_{1-4})alkyl, 4-butanolidyl, 5-pentanolidyl, tetrahydrothiopyranyl, 1-oxotetrahydrothiopyranyl, 1,1-dioxotetrahydrothiopyranyl, C_{1-4} alkyl [substituted by 1 or 2 R^8 groups (provided that when there are 2 R^8 groups they are not substituents on the same carbon)] and groups of the formulae B and B':



(B)



wherein y is 0 or 1, t is 0, 1, 2 or 3 and u is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen);

{ wherein R^8 is independently selected from hydroxy, C_{1-4} alkoxy, C_{1-4} alkoxy, hydroxy C_{1-4} alkoxy, 2,2-dimethyl-1,3-dioxolan-4-yl, heterocyclyl, C_{1-4} alkanoyl, C_{1-4} alkoxy, C_{1-4} alkanesulfinyl, C_{1-4} alkanesulfonyl, $-N(OH)CHO$, $-COCOOR^9$, $(R^9)(R^{10})NCO-$, $(R^9)(R^{10})NSO_2-$, $-COCH_2OR^{11}$, $(R^9)(R^{10})N-$ and $-COOR^9$;

[wherein R^9 and R^{10} are independently selected from hydrogen, hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), C_{5-7} cycloalkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano(C_{1-4})alkyl, 4-butanolidyl, 5-pentanolidyl, tetrahydrothiopyranyl, 1-oxotetrahydrothiopyranyl, 1,1-dioxotetrahydrothiopyranyl, 2,2-dimethyl-1,3-dioxolan-4-yl and C_{1-4} alkyl substituted by R^{13} ;

(wherein R^{13} is selected from hydroxy, C_{1-4} alkoxy, heterocyclyl, C_{1-4} alkanoyl, C_{1-4} alkanesulfinyl, C_{1-4} alkanesulfonyl, $-N(OH)CHO$, $(R^{11})(R^{12})NCO-$, $(R^{11})(R^{12})NSO_2-$, $-COCH_2OR^{11}$, $(R^{11})(R^{12})N-$

{ wherein R^{11} and R^{12} are independently selected from hydrogen, C_{1-4} alkyl, C_{1-4} alkoxy, hydroxy C_{1-4} alkyl, C_{1-4} alkylS(O)_b (wherein b is 0, 1 or 2)); and

R⁹ and R¹⁰ can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from oxo, hydroxy, carboxy, halo, nitro, nitroso, cyano, isocyano, amino, *N*-C₁₋₄alkylamino, *N,N*-(C₁₋₄)₂alkylamino, carbonyl, sulfo, C₁₋₄alkoxy, heterocyclyl, C₁₋₄alkanoyl, C₁₋₄alkanesulfinyl, C₁₋₄alkanesulfonyl, -N(OH)CHO, (R¹¹)(R¹²)NCO-, (R¹¹)(R¹²)NSO₂-, -COCH₂OR¹¹, (R¹¹)(R¹²)N-; wherein R¹¹ and R¹² are as defined above] };

provided that when R¹ is of the formula A' or A'' then R³ does not contain a group of the formula B or B' and when R³ is of the formula B or B' then R¹ does not contain a group of the formula A' or A'';

or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof;

with the proviso that the compound of formula (1) is not:

- i) 2,3-dichloro-5-[*N*-(2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-4*H*-thieno[3,2-*b*]pyrrole;
- ii) 2-chloro-5-[*N*-(2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6*H*-thieno[2,3-*b*]pyrrole; or
- iii) 2-chloro-5-[*N*-(1-methyl-2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6*H*-thieno[2,3-*b*]pyrrole.

In another aspect, the invention relates to compounds of formula (1) as hereinabove defined or to a pharmaceutically acceptable salt.

It is to be understood that, insofar as certain of the compounds of formula (1) defined above may exist in optically active or racemic forms by virtue of one or more asymmetric carbon atoms, the invention includes in its definition any such optically active or racemic form which possesses glycogen phosphorylase inhibition activity. The synthesis of optically active forms may be carried out by standard techniques of organic chemistry well known in the art, for example by synthesis from optically active starting materials or by resolution of a racemic form. Similarly, the above-mentioned activity may be evaluated using the standard laboratory techniques referred to hereinafter.

Within the present invention it is to be understood that a compound of the formula (1) or a salt thereof may exhibit the phenomenon of tautomerism and that the formulae drawings within this specification can represent only one of the possible

tautomeric forms. It is to be understood that the invention encompasses any tautomeric form which has glycogen phosphorylase inhibition activity and is not to be limited merely to any one tautomeric form utilised within the formulae drawings. The formulae drawings within this specification can represent only one of the possible tautomeric forms and it is to be understood that the specification encompasses all possible tautomeric forms of the compounds drawn not just those forms which it has been possible to show graphically herein.

It is also to be understood that certain compounds of the formula (1) and salts thereof can exist in solvated as well as unsolvated forms such as, for example, hydrated forms. It is to be understood that the invention encompasses all such solvated forms which have glycogen phosphorylase inhibition activity.

It is also to be understood that certain compounds of the formula (1) may exhibit polymorphism, and that the invention encompasses all such forms which possess glycogen phosphorylase inhibition activity.

The present invention relates to the compounds of formula (1) as hereinbefore defined as well as to the salts thereof. Salts for use in pharmaceutical compositions will be pharmaceutically acceptable salts, but other salts may be useful in the production of the compounds of formula (1) and their pharmaceutically acceptable salts. Pharmaceutically acceptable salts of the invention may, for example, include acid addition salts of the compounds of formula (1) as hereinbefore defined which are sufficiently basic to form such salts. Such acid addition salts include for example salts with inorganic or organic acids affording pharmaceutically acceptable anions such as with hydrogen halides (especially hydrochloric or hydrobromic acid of which hydrochloric acid is particularly preferred) or with sulphuric or phosphoric acid, or with trifluoroacetic, citric or maleic acid. Suitable salts include hydrochlorides, hydrobromides, phosphates, sulphates, hydrogen sulphates, alkylsulphonates, arylsulphonates, acetates, benzoates, citrates, maleates, fumarates, succinates, lactates and tartrates. In addition where the compounds of formula (1) are sufficiently acidic, pharmaceutically acceptable salts may be formed with an inorganic or organic base which affords a pharmaceutically acceptable cation. Such salts with inorganic or organic bases include for example an alkali metal salt, such as a sodium or potassium salt, an alkaline earth metal salt such as a calcium or magnesium salt, an ammonium salt or for example a

salt with methylamine, dimethylamine, trimethylamine, piperidine, morpholine or tris-(2-hydroxyethyl)amine.

An *in vivo* hydrolysable ester of a compound of formula (1) containing carboxy or hydroxy group is, for example. A pharmaceutically acceptable ester which is cleaved in the human or animal body to produce the parent acid or alcohol.

Suitable pharmaceutically acceptable esters for carboxy include C₁₋₆alkoxymethyl esters for example methoxymethyl, C₁₋₆alkanoyloxymethyl esters for example pivaloyloxymethyl, phthalidyl esters, C₃₋₈cycloalkoxycarbonyloxyC₁₋₆alkyl esters for example 1-cyclohexylcarbonyloxyethyl; 1,3-dioxolen-2-onylmethyl esters for example 5-methyl-1,3-dioxolen-2-onylmethyl; and C₁₋₆alkoxycarbonyloxyethyl esters for example 1-methoxycarbonyloxyethyl and may be formed at any carboxy group in the compounds of this invention.

Suitable pharmaceutically-acceptable esters for hydroxy include inorganic esters such as phosphate esters (including phosphoramidic cyclic esters) and α -acyloxyalkyl ethers and related compounds which as a result of the *in-vivo* hydrolysis of the ester breakdown to give the parent hydroxy group/s. Examples of α -acyloxyalkyl ethers include acetoxymethoxy and 2,2-dimethylpropionyloxymethoxy. A selection of *in-vivo* hydrolysable ester forming groups for hydroxy include C₁₋₁₀alkanoyl, for example acetyl; benzoyl; phenylacetyl; substituted benzoyl and phenylacetyl, C₁₋₁₀alkoxycarbonyl (to give alkyl carbonate esters), for example ethoxycarbonyl; di-(C₁₋₄)alkylcarbamoyl and *N*-(di-(C₁₋₄)alkylaminoethyl)-*N*-(C₁₋₄)alkylcarbamoyl (to give carbamates); di-(C₁₋₄)alkylaminoacetyl and carboxyacetyl. Examples of ring substituents on phenylacetyl and benzoyl include aminomethyl, (C₁₋₄)alkylaminomethyl and di-((C₁₋₄)alkyl)aminomethyl, and morpholino or piperazino linked from a ring nitrogen atom via a methylene linking group to the 3- or 4- position of the benzoyl ring. Other interesting *in-vivo* hydrolysable esters include, for example, R^AC(O)O(C₁₋₆)alkyl-CO-, wherein R^A is for example, benzyloxy-(C₁₋₄)alkyl, or phenyl). Suitable substituents on a phenyl group in such esters include, for example, 4-(C₁₋₄)piperazino-(C₁₋₄)alkyl, piperazino-(C₁₋₄)alkyl and morpholino-(C₁₋₄)alkyl.

In this specification the generic term "alkyl" includes both straight-chain and branched-chain alkyl groups. However references to individual alkyl groups such as "propyl" are specific for the straight chain version only and references to individual branched-chain alkyl groups such as *t*-butyl are specific for the branched chain version only. For example, "C₁₋₄alkyl" includes methyl, ethyl, propyl, isopropyl and *t*-butyl and examples of "C₁₋₆alkyl" include the examples of "C₁₋₄alkyl" and additionally pentyl, 2,3-dimethylpropyl, 3-methylbutyl and hexyl. An analogous convention applies to other generic terms, for example "C₂₋₄alkenyl" includes vinyl, allyl and 1-propenyl and examples of "C₂₋₆alkenyl" include the examples of "C₂₋₄alkenyl" and additionally 1-butenyl, 2-butenyl, 3-butenyl, 2-methylbut-2-enyl, 3-methylbut-1-enyl, 1-pentenyl, 3-pentenyl and 4-hexenyl. Examples of "C₂₋₄alkynyl" includes ethynyl, 1-propynyl and 2-propynyl and examples of "C₂₋₆alkynyl" include the examples of "C₂₋₄alkynyl" and additionally 3-butyne, 2-pentyne and 1-methylpent-2-ynyl.

The term "hydroxyC₁₋₄alkyl" includes hydroxymethyl, hydroxyethyl, hydroxypropyl, hydroxyisopropyl and hydroxybutyl. The term "hydroxyethyl" includes 1-hydroxyethyl and 2-hydroxyethyl. The term "hydroxypropyl" includes 1-hydroxypropyl, 2-hydroxypropyl and 3-hydroxypropyl and an analogous convention applies to terms such as hydroxybutyl. The term "dihydroxyC₁₋₄alkyl" includes dihydroxymethyl, dihydroxyethyl, dihydroxypropyl, dihydroxyisopropyl and dihydroxybutyl. The term "dihydroxyethyl" includes 1,1-dihydroxyethyl 2,2-dihydroxyethyl and 1,2-dihydroxyethyl. An analogous convention applies to terms such as dihydroxypropyl, dihydroxyisopropyl and dihydroxybutyl.

The term "halo" refers to fluoro, chloro, bromo and iodo.

Examples of "C₁₋₄alkoxy" include methoxy, ethoxy, propoxy and isopropoxy. Examples of "C₁₋₆alkoxy" include the examples of "C₁₋₄alkoxy" and additionally butyloxy, *t*-butyloxy, pentoxy and 1,2-(methyl)₂propoxy. Examples of "C₁₋₄alkanoyl" include formyl, acetyl and propionyl. Examples of "C₁₋₆alkanoyl" include the example of "C₁₋₄alkanoyl" and additionally butanoyl, pentanoyl, hexanoyl and 1,2-(methyl)₂propionyl. Examples of "C₁₋₄alkanoyloxy" are formyloxy, acetoxy and propionyloxy. Examples of "C₁₋₆alkanoyloxy" include the examples of "C₁₋₄alkanoyloxy" and additionally butanoyloxy, pentanoyloxy, hexanoyloxy and 1,2-(methyl)₂propionyloxy. Examples of "*N*-(C₁₋₄alkyl)amino" include methylamino and ethylamino. Examples of "*N*-(C₁₋₆alkyl)amino" include the examples of

"*N*-(C_{1-4} alkyl)amino" and additionally pentylamino, hexylamino and 3-methylbutylamino. Examples of "*N,N*-(C_{1-4} alkyl)₂amino" include *N,N*-(methyl)₂amino, *N,N*-(ethyl)₂amino and *N*-ethyl-*N*-methylamino. Examples of "*N,N*-(C_{1-6} alkyl)₂amino" include the example of "*N,N*-(C_{1-4} alkyl)₂amino" and additionally *N*-methyl-*N*-pentylamino and *N,N*-(pentyl)₂amino. Examples of "*N*-(C_{1-4} alkyl)carbamoyl" are methylcarbamoyl and ethylcarbamoyl. Examples of "*N*-(C_{1-6} alkyl)carbamoyl" are the examples of "*N*-(C_{1-4} alkyl)carbamoyl" and additionally pentylcarbamoyl, hexylcarbamoyl and 1,2-(methyl)₂propylcarbamoyl. Examples of "*N,N*-(C_{1-4} alkyl)₂carbamoyl" are *N,N*-(methyl)₂carbamoyl, *N,N*-(ethyl)₂carbamoyl and *N*-methyl-*N*-ethylcarbamoyl. Examples of "*N,N*-(C_{1-6} alkyl)₂carbamoyl" are the examples of "*N,N*-(C_{1-4} alkyl)₂carbamoyl" and additionally *N,N*-(pentyl)₂carbamoyl, *N*-methyl-*N*-pentylcarbamoyl and *N*-ethyl-*N*-hexylcarbamoyl. Examples of "*N*-(C_{1-4} alkyl)sulphamoyl" are *N*-(methyl)sulphamoyl and *N*-(ethyl)sulphamoyl. Examples of "*N*-(C_{1-6} alkyl)sulphamoyl" are the examples of "*N*-(C_{1-4} alkyl)sulphamoyl" and additionally *N*-pentylsulphamoyl, *N*-hexylsulphamoyl and 1,2-(methyl)₂propylsulphamoyl. Examples of "*N,N*-(C_{1-4} alkyl)₂sulphamoyl" are *N,N*-(methyl)₂sulphamoyl, *N,N*-(ethyl)₂sulphamoyl and *N*-(methyl)-*N*-(ethyl)sulphamoyl. Examples of "*N,N*-(C_{1-6} alkyl)₂sulphamoyl" are the examples of "*N,N*-(C_{1-4} alkyl)₂sulphamoyl" and additionally *N,N*-(pentyl)₂sulphamoyl, *N*-methyl-*N*-pentylsulphamoyl and *N*-ethyl-*N*-hexylsulphamoyl.

Examples of "cyano(C_{1-4} alkyl)" are cyanomethyl, cyanoethyl and cyanopropyl. Examples of " C_{5-7} cycloalkyl" are cyclopentyl, cyclohexyl and cycloheptyl. Examples of " C_{3-8} cycloalkyl" include " C_{5-7} cycloalkyl, cyclopropyl, cyclobutyl and cyclooctyl.

The term "amino C_{1-4} alkyl" includes aminomethyl, aminoethyl, aminopropyl, aminoisopropyl and aminobutyl. The term "aminoethyl" includes 1-aminoethyl and 2-aminoethyl. The term "aminopropyl" includes 1-aminopropyl, 2-aminopropyl and 3-aminopropyl and an analogous convention applies to terms such as aminoethyl and aminobutyl.

Examples of " C_{1-6} alkylS(O)_a (wherein a is 0 to 2)" include methylthio, ethylthio, propylthio, methanesulphinyl, ethanesulphinyl, propanesulphinyl, mesyl, ethanesulphonyl, propanesulphonyl, isopropanesulphonyl, pentanesulphonyl and hexanesulphonyl.

Examples of " C_{1-4} alkylS(O)_c (wherein c is 0 to 2)", " C_{1-4} alkylS(O)_d (wherein d is 0 to 2)", " C_{1-4} alkylS(O)_e (wherein e is 0 to 2)", and " C_{1-4} alkylS(O)_f (wherein f is 0 to 2)"

independently include methylthio, ethylthio, propylthio, methanesulphinyl, ethanesulphinyl, propanesulphinyl, mesyl, ethanesulphonyl, propanesulphonyl and isopropanesulphonyl.

Examples of "C₁₋₆alkoxycarbonyl" include methoxycarbonyl, ethoxycarbonyl, *n*- and *t*-butoxycarbonyl. Examples of "C₁₋₆alkoxycarbonylamino" include methoxycarbonylamino, ethoxycarbonylamino, *n*- and *t*-butoxycarbonylamino. Examples of "C₁₋₆alkylsulphonyl-*N*-(C₁₋₆alkyl)amino" include methylsulphonyl-*N*-methyldamino, ethylsulphonyl-*N*-methyldamino and propylsulphonyl-*N*-ethylamino. Examples of "C₁₋₆alkylsulphonylamino" include methylsulphonylamino, ethylsulphonylamino and propylsulphonylamino. Examples of "C₁₋₆alkanoylamino" include formamido, acetamido and propionylamino.

Examples of "C₁₋₄alkoxyC₁₋₄alkoxy" are methoxymethoxy, ethoxymethoxy, ethoxyethoxy and methoxyethoxy. Examples of "hydroxyC₁₋₄alkoxy" are hydroxyethoxy and hydroxypropoxy. Examples of "hydroxypropoxy" are 1-hydroxypropoxy, 2-hydroxypropoxy and 3-hydroxypropoxy.

Where optional substituents are chosen from "0, 1, 2 or 3" groups it is to be understood that this definition includes all substituents being chosen from one of the specified groups or the substituents being chosen from two or more of the specified groups. An analogous convention applies to substituents chose from "0, 1 or 2" groups and "1 or 2" groups.

"Heterocyclyl" is a saturated, partially saturated or unsaturated, monocyclic ring containing 5 to 7 atoms of which 1, 2, 3 or 4 ring atoms are chosen from nitrogen, sulphur or oxygen, which may, unless otherwise specified, be carbon or nitrogen linked, wherein a -CH₂- group can optionally be replaced by a -C(O)- and a ring sulphur atom may be optionally oxidised to form the S-oxide(s). Examples and suitable values of the term "heterocyclyl" are morpholino, morpholinyl, piperidino, piperidyl, pyridyl, pyranyl, pyrrolyl, imidazolyl, thiazolyl, thienyl, dioxolanyl, thiadiazolyl, piperazinyl, isothiazolidinyl, triazolyl, tetrazolyl, pyrrolidinyl, 2-oxazolidinonyl, 5-isoxazolonyl, thiomorpholino, pyrrolinyl, homopiperazinyl, 3,5-dioxapiperidinyl, 3-oxopyrazolin-5-yl, tetrahydropyranyl, , pyrimidyl, pyrazinyl, pyridazinyl, isoxazolyl, 4-oxopyridyl, 2-oxopyrrolidyl, 4-oxothiazolidyl, furyl, thienyl, oxazolyl, and oxadiazolyl. Preferably a "heterocyclyl" is morpholino, morpholinyl, piperidino, piperidyl, pyridyl, pyranyl, pyrrolyl, imidazolyl, thiazolyl, thienyl, thiadiazolyl, piperazinyl, isothiazolidinyl,

...1,3,4-triazolyl, tetrazolyl, pyrrolidinyl, thiomorpholino, pyrrolinyl, homopiperazinyl, 3,5-dioxapiperidinyl, pyrimidyl, pyrazinyl, pyridazinyl, isoxazolyl, 4-oxopyridyl, 2-oxopyrrolidyl, 4-oxothiazolidyl, furyl, thienyl, oxazolyl and 1,2,4-oxadiazolyl. More preferably heterocyclyl is oxazolyl, 1,2,4-oxadiazolyl, pyridyl, furyl, thienyl, morpholino, pyrazinyl and piperazinyl.

Examples of "(heterocyclyl) C_{1-4} alkyl" are morpholinomethyl, morpholinethyl, morpholinylmethyl, morpholinylethyl, piperidinomethyl, piperidinoethyl, piperidylmethyl, piperidylethyl, imidazolylmethyl, imidazolylethyl, oxazolylmethyl, oxazolylethyl 1,2,4-oxadiazolylmethyl, 1,2,4-oxadiazolylethyl, pyridylmethyl, pyridylethyl, furylmethyl, furylethyl, (thienyl)methyl, (thienyl)ethyl, pyrazinylmethyl, pyrazinylethyl, piperazinylmethyl and piperazinylethyl.

Examples of "aryl" are phenyl and naphthyl.

Examples of "aryl(C_{1-4})alkyl" are benzyl, phenethyl, naphthylmethyl and naphthylethyl.

"Heteroarylene" is a diradical of a heteroaryl group. A heteroaryl group is aryl, monocyclic ring containing 5 to 7 atoms of which 1, 2, 3 or 4 ring atoms are chosen from nitrogen, sulphur or oxygen. Examples of heteroarylene are pyridylene, pyrimidinylene, pyrazinylene, pyridazinylene, pyrrolylene, thienylene and furylene.

Preferred values of A, R^1 , R^2 , R^3 , R^4 , R^5 and n are as follows. Such values may be used where appropriate with any of the definitions, claims or embodiments defined hereinbefore or hereinafter.

In one aspect of the present invention there is provided a compound of formula (1) as depicted above wherein R^4 and R^5 are together $-S-C(R^6)=C(R^7)-$.

In another aspect of the invention R^4 and R^5 are together $-C(R^7)=C(R^6)-S-$.

In a further aspect of the invention, R^6 and R^7 are independently selected from hydrogen, halo or C_{1-6} alkyl.

Preferably R^6 and R^7 are independently selected from hydrogen, chloro, bromo or methyl.

Particularly R^6 and R^7 are independently selected from hydrogen or chloro.

More particularly one of R^6 and R^7 is chloro.

In one aspect of the invention A is phenylene.

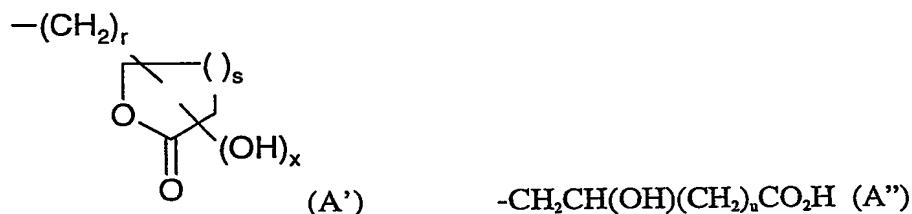
In another aspect of the invention A is heteroarylene.

5 Preferably A is selected from phenylene, pyridylene, pyrimidinylene, pyrrolylene, thienylene and furylene.

In one aspect of the invention n is 0 or 1.

Preferably n is 1.

10 In another aspect of the present invention R^1 is selected from hydrogen, halo, nitro, cyano, hydroxy, fluoromethyl, difluoromethyl, trifluoromethyl, C_{1-4} alkoxy and R^1 is of the formula A' or A'':



15 wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

Preferably R^1 is hydrogen or halo.

More preferably R^1 is hydrogen.

In one aspect of the invention s is 1.

20 In another aspect of the invention s is 2.

In one aspect of the invention --- is a single bond.

In another aspect of the invention = is a double bond.

25 In one aspect of the invention R^2 is hydrogen.

In another aspect of the invention R^2 is carboxy.

In another aspect of the invention R^2 is hydroxy.

In one aspect on the invention R^3 is selected from hydrogen, hydroxy; C_{1-4} alkanoyl, carbamoyl, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), C_{5-7} cycloalkyl (optionally substituted with 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano(C_{1-4})alkyl, 4-butanolidyl, 5-pentanolidyl, tetrahydrothiopyranyl, 1-oxotetrahydrothiopyranyl, 1,1-dioxotetrahydrothiopyranyl and C_{1-4} alkyl [substituted by 1 or 2 R^8 (provided that when there are 2 R^8 groups they are not substituents on the same carbon)];

{ wherein R^8 is independently selected from hydroxy, C_{1-4} alkoxy, C_{1-4} alkoxy, hydroxy C_{1-4} alkoxy, 2,2-dimethyl-1,3-dioxolan-4-yl, 1,2,4-oxadiazolyl, tetrazolyl, imidazolyl, pyrrolidinyl, piperidyl, tetrahydrofuryl, tetrahydropyranyl, tetrahydrothiopyranyl group, tetrahydrothienyl, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanesulfinyl, C_{1-4} alkanesulfonyl, $-N(OH)CHO$, $-COCOOR^9$, $(R^9)(R^{10})NCO-$, $-COCH_2OR^{11}$, $(R^9)(R^{10})N-$ and $-COOR^9$;

[wherein R^9 and R^{10} are independently selected from hydrogen, hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), C_{5-7} cycloalkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano(C_{1-4})alkyl, 4-butanolidyl, 5-pentanolidyl, tetrahydrothiopyranyl, 1-oxotetrahydrothiopyranyl, 1,1-dioxotetrahydrothiopyranyl, 2,2-dimethyl-1,3-dioxolan-4-yl and C_{1-4} alkyl substituted by R^{13} ;

(wherein R^{13} is selected from hydroxy, C_{1-4} alkoxy, 1,2,4-oxadiazolyl, tetrazolyl, imidazolyl, pyrrolidinyl, piperidyl, tetrahydrofuryl, tetrahydropyranyl, tetrahydrothiopyranyl, tetrahydrothienyl, C_{1-4} alkanoyl, C_{1-4} alkanesulfinyl, C_{1-4} alkanesulfonyl, $-N(OH)CHO$, $(R^{11})(R^{12})NCO-$, $(R^{11})(R^{12})NSO_2-$, $-COCH_2OR^{11}$, $(R^{11})(R^{12})N-$;

{ wherein R^{11} and R^{12} are independently selected from hydrogen, C_{1-4} alkyl, C_{1-4} alkoxy, hydroxy C_{1-4} alkyl, C_{1-4} alkylS(O)_b (wherein b is 0, 1 or 2)); and

R^9 and R^{10} can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from oxo, hydroxy, carboxy, halo, nitro, nitroso, cyano, isocyano, amino, N - C_{1-4} alkylamino, N,N -(C_{1-4})₂alkylamino, carbonyl, sulfo, C_{1-4} alkoxy, heterocyclyl, C_{1-4} alkanoyl, C_{1-4} alkanesulfinyl, C_{1-4} alkanesulfonyl, $-N(OH)CHO$, $(R^{11})(R^{12})NCO-$,

$(R^{11})(R^{12})NSO_2$, $-COCH_2OR^{11}$, $(R^{11})(R^{12})N$ - and wherein R^{11} and R^{12} are as defined above]];

In a further aspect of the invention R^3 is selected from C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano C_{1-4} alkyl and C_{1-4} alkyl [substituted by 1 or 2 of R^8 groups (provided that when there are 2 R^8 groups they are not substituents on the same carbon)];

{ wherein R^8 is independently selected from hydroxy, C_{1-4} alkoxy C_{1-4} alkoxy, hydroxy C_{1-4} alkoxy, 2,2-dimethyl-1,3-dioxolan-4-yl, heterocyclyl, C_{1-4} alkoxy, C_{1-4} alkanesulfinyl, C_{1-4} alkanesulfonyl, $(R^9)(R^{10})NCO$ -, $-COCOOR^9$, $-COOR^9$, $(R^9)(R^{10})N$ - ;

[wherein R^9 and R^{10} are independently selected from hydrogen, hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C_{1-4} alkyl substituted with R^{13} (wherein R^{13} is C_{1-4} alkoxy); and

R^9 and R^{10} can together with the nitrogen to which they are attached form a 4- to 6-membered ring where the ring may be optionally substituted on carbon by 1 or 2 hydroxy groups or carboxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon)]}.

In yet a further aspect of the inventions R^3 is selected from hydroxyethyl, hydroxypropyl, 1,3-dihydroxyprop-2-yl, 3-hydroxy-2-hydroxymethyl-propyl, 2,3-dihydroxypropyl, 2,2-dimethyl-1,3-dioxolan-4-ylmethyl, cyanomethyl, cyanoethyl, cyanopropyl, carbamoyl, carboxycarbonyl, methanoyl, ethanoyl, propanoyl, methoxymethyl, ethoxymethyl, methoxyethyl, ethoxymethyl, (2,3-dihydroxypyrrolidinyl)carbonylmethyl, (3-hydroxypiperidino)carbonylmethyl, 3-carboxyazolidin-1-ylcarbonylmethyl, 1,2,4-oxadiazolylmethyl, tetrazolylmethyl, imidazolylmethyl, pyrrolidinylmethyl, piperidylmethyl, tetrahydrofurylmethyl, tetrahydropyranylmethyl, (tetrahydrothiopyranyl)methyl, (tetrahydrothienyl)methyl, methanesulfinylethyl, methanesulfonylethyl, *N*-(1,3-dihydroxyprop-2-yl)carbamoylmethyl, 2-(methylamino)ethyl, 2-(dimethylamino)ethyl, 2-(ethylamino)ethyl, 2-(diethylamino)ethyl, 2-(*N*-methyl-*N*-ethylamino)ethyl, methoxycarbonylmethyl, carboxymethyl, carbomoylmethyl, (dimethylcarbamoyl)methyl,

(methylcarbamoyl)methyl; (hydroxycabamoyl)methyl, (hydroxyethylcarbamoyl)methyl, (methoxyethylcarbamoyl)methyl and 2-(4-hydroxypiperidino)-2-oxoethyl.

*A preferred class of compound is of the formula (1) wherein;

5 --- is a single bond;

X is CH;

R^4 and R^5 are together $-\text{C}(R^7)=\text{C}(R^6)-\text{S}-$;

R^6 is halo;

10

R^7 is hydrogen;

A is phenylene;

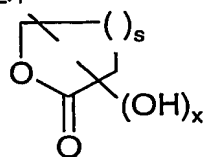
15

n is 1 or 2;

R^1 is independently selected from hydrogen, halo, cyano, nitro, hydroxy, fluoromethyl, difluoromethyl, trifluoromethyl, C_{1-4} alkoxy and

R^1 is of the formula A' or A'':

$-(\text{CH}_2)_r$



(A')

$-\text{CH}_2\text{CH}(\text{OH})(\text{CH}_2)_n\text{CO}_2\text{H}$ (A'')

20

wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

R^2 is hydrogen;

25

R^3 is selected from C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano C_{1-4} alkyl, and C_{1-4} alkyl [substituted by 1 or 2 R^8 groups (provided that when there are 2 R^8 groups they are not substituents on the same carbon)];

{R⁸ is independently selected from hydroxy, heterocyclyl, C₁₋₄alkanoyl, C₁₋₄alkoxy, C₁₋₄alkanesulfinyl, C₁₋₄alkanesulfonyl, -COCOOR⁹, (R⁹)(R¹⁰)NCO-, -COCH₂OR¹¹, (R⁹)(R¹⁰)N-, -COOR⁹ and 2,2-dimethyl-1,3-dioxolan-4-yl;

[R⁹ and R¹⁰ are independently selected from hydrogen, hydroxy, C₁₋₄alkyl

- 5 (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C₁₋₄alkyl substituted by C₁₋₄alkoxy and wherein R⁹ and R¹⁰ can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

- 10 R¹¹ is selected from hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy and hydroxyC₁₋₄alkyl}};
or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof;
with the proviso that the compound of formula (1) is not:

- 15 i. 2-chloro-5-[N-(2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6*H*-thieno[2,3-*b*]pyrrole; or
ii. 2-chloro-5-[N-(1-methyl-2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6*H*-thieno[2,3-*b*]pyrrole.

Another preferred class of compounds is of formula (1) wherein:

- 20 --- is a single bond;

X is CH;

R⁴ and R⁵ are together -C(R⁷)=C(R⁶)-S-;

R⁶ is chloro;

25

R⁷ is hydrogen;

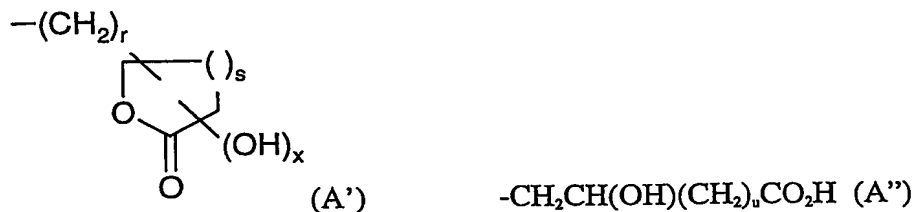
A is phenylene;

30

n is 1 or 2;

4alkoxy and

R^1 is of the formula A' or A'' :



- 5 wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not
a substituent on the ring carbon adjacent to the ring oxygen;

R² is hydrogen;

- 10 R³ is selected from C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyanoC₁₋₄alkyl, and C₁₋₄alkyl [substituted by 1 or 2 R⁸ groups (provided that when there are 2 R⁸ groups they are not substituents on the same carbon)];

{R⁸ is independently selected from hydroxy, heterocyclyl, C₁₋₄alkanoyl, C₁₋₄alkoxy, C₁₋₄alkanesulfinyl, C₁₋₄alkylsulfonyl, -COCOOR⁹, (R⁹)(R¹⁰)NCO-, -COCH₂OR¹¹, (R⁹)(R¹⁰)N-, -COOR⁹ and 2,2-dimethyl-1,3-dioxolan-4-yl;

[R⁹ and R¹⁰ are independently selected from hydrogen, hydroxy, C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C₁₋₄alkyl substituted by C₁₋₄alkoxy and wherein R⁹ and R¹⁰ can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

R¹¹ is selected from hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy and hydroxyC₁₋₄alkyl}};
or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof;

- 25 with the proviso that the compound of formula (1) is not:

- i. 2-chloro-5-[*N*-(2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6*H*-thieno[2,3-*b*]pyrrole; or

- ii. 2-chloro-5-[N-(1-methyl-2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6H-thieno[2,3-b]pyrrole.

5 Another preferred class of compound is of the formula (1) wherein:

----- is a single bond;

X is CH;

R⁴ and R⁵ are together -S-C(R⁶)=C(R⁷)-;

10 R⁶ is hydrogen or halo;

R⁷ is hydrogen or halo;

A is phenylene;

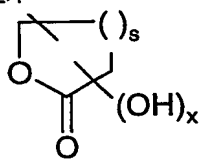
15

n is 1 or 2;

R¹ is independently selected from hydrogen, halo, nitro, hydroxy, C₁₋₄alkyl, C₁₋₄alkoxy and

20 R¹ is of the formula A' or A'':

-(CH₂)_r



(A')

-CH₂CH(OH)(CH₂)_nCO₂H (A'')

wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

25

R² is hydrogen;

R³ is selected from C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same

carbon); cyanoC₁₋₄alkyl; and C₁₋₄alkyl [substituted by 1 or 2-R⁸ groups (provided that when there are 2 R⁸ groups they are not substituents on the same carbon)];

{R⁸ is independently selected from hydroxy, heterocyclyl, C₁₋₄alkanoyl, C₁₋₄alkoxy, C₁₋₄alkanesulfinyl, C₁₋₄alkylsulfonyl, -COCOOR⁹, (R⁹)(R¹⁰)NCO-, -

5 COCH₂OR¹¹, (R⁹)(R¹⁰)N-, -COOR⁹ and 2,2-dimethyl-1,3-dioxolan-4-yl;

[R⁹ and R¹⁰ are independently selected from hydrogen, hydroxy, C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C₁₋₄alkyl substituted by C₁₋₄alkoxy and wherein R⁹ and R¹⁰ can together with the nitrogen to which they are attached
10 form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

R¹¹ is selected from hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy and hydroxyC₁₋₄alkyl];
or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof;

15 with the proviso that the compound of formula (1) is not 2,3-dichloro-5-[N-(2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-4H-thieno[3,2-b]pyrrole.

A further preferred class of compound is of the formula (1) wherein;

---- is a single bond;

X is CH;

20 R⁴ and R⁵ are together -C(R⁷)=C(R⁶)-S-;

R⁶ is halo;

R⁷ is hydrogen;

25

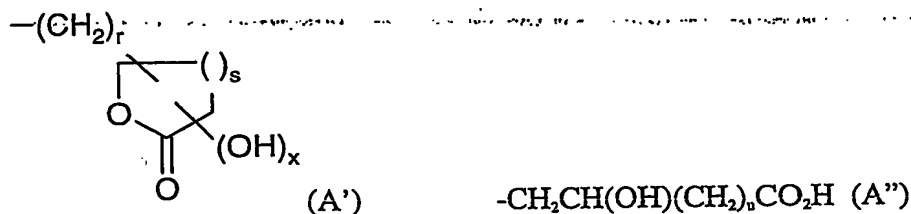
A is phenylene;

n is 1 or 2;

30

R¹ is independently selected from hydrogen, halo, nitro, hydroxy, C₁₋₄alkyl, C₁₋₄alkoxy and

R¹ is of the formula A' or A":



wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

5 R^2 is hydrogen;

R^3 is selected from C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon);

10 or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof; with the proviso that the compound of formula (1) is not:

- i. 2-chloro-5-[N-(2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6*H*-thieno[2,3-*b*]pyrrole; or
- 15 ii. 2-chloro-5-[N-(1-methyl-2-oxo-1,2,3,4-tetrahydroquinol-3-yl)carbamoyl]-6*H*-thieno[2,3-*b*]pyrrole.

A further preferred class of compound is of the formula (1) wherein;

20 --- is a single bond;

X is CH;

R^4 and R^5 are together $-C(R^7)=C(R^6)-S-$;

R^6 is halo;

25

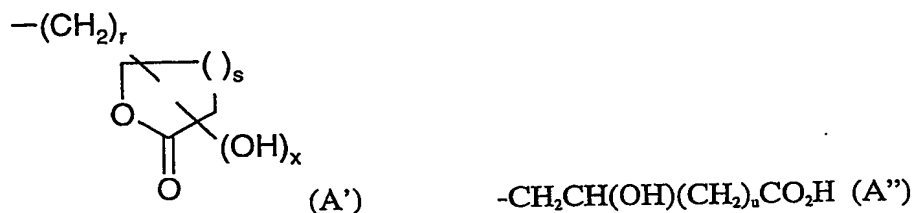
R^7 is hydrogen;

A is phenylene;

n is 1 or 2;

R¹ is independently selected from hydrogen, halo, nitro, hydroxy, C₁₋₄alkyl, C₁₋₄alkoxy and

5 R^1 is of the formula A' or A'' :



wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

10 R^2 is hydrogen;

R³ is selected from cyanoC₁₋₄alkyl, and C₁₋₄alkyl substituted by R⁸;

R^8 is selected from $(R^9)(R^{10})NCO-$, $(R^9)(R^{10})N-$, and $-COOR^9$;

R⁹ and R¹⁰ are independently selected from hydrogen, hydroxy, C₁₋₄alkyl

15 (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C₁₋₄alkyl substituted by C₁₋₄alkoxy and wherein R⁹ and R¹⁰ can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

20 or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof.

Another class of compounds is of the formula (1) wherein

== is a double bond;

X is CH;

25 R⁴ and R⁵ are together -C(R⁷)=C(R⁶)-S- or -S-C(R⁷)=C(R⁶)-;

R^6 is hydrogen or halo;

R^7 is hydrogen or hydrogen;

or a pharmaceutically acceptable salt or *in-vivo* hydrolysable ester thereof.

A further class of compound is of formula (1) wherein:

— is a single bond;

5 X is CH;

R⁴ and R⁵ are together —C(R⁷)=C(R⁶)-S- or —S-C(R⁷)=C(R⁶)-;

R⁶ is hydrogen or halo;

10 R⁷ is hydrogen or hydrogen;

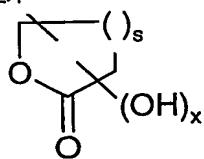
A is pyridylene;

n is 1 or 2;

15

R¹ is independently selected from hydrogen, halo, nitro, hydroxy, C₁₋₄alkyl, C₁₋₄alkoxy and R¹ is of the formula A' or A'':

—(CH₂)_r



(A')

—CH₂CH(OH)(CH₂)_uCO₂H (A'')

wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not
20 a substituent on the ring carbon adjacent to the ring oxygen;

R² is hydrogen;

25 R³ is selected from hydroxy, C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyanoC₁₋₄alkyl, and C₁₋₄alkyl [substituted by 1 or 2 R⁸ (provided that when there are 2 R⁸ groups they are not substituents on the same carbon)];

{R⁸ is selected from hydroxy, C₁₋₄alkoxyC₁₋₄alkoxy, hydroxyC₁₋₄alkoxy, heterocyclyl, C₁₋₄alkanoyl, C₁₋₄alkoxy, C₁₋₄alkanesulfinyl, C₁₋₄alkylsulfonyl, —COCOOR⁹,

(R⁹)(R¹⁰)NCO-, -COCH₂OR¹¹, (R⁹)(R¹⁰)N- γ -COOR⁹ and 2,2-dimethyl-1,3-dioxolan-4-yl;

[R⁹ and R¹⁰ are independently selected from hydrogen, hydroxy, C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C₁₋₄alkyl substituted by C₁₋₄alkoxy and wherein R⁹ and R¹⁰ can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

R¹¹ is selected from hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy and hydroxyC₁₋₄alkyl];

or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof.

Yet another preferred class of compound is of the formula (1) wherein;

----- is a single bond;

X is N;

R⁴ and R⁵ are together -C(R⁷)=C(R⁶)-S-;

R⁶ is halo;

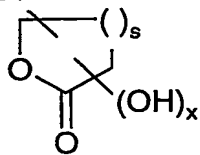
R⁷ is hydrogen;

A is phenylene;

n is 1 or 2;

R¹ is independently selected from hydrogen, halo, cyano, nitro, hydroxy, fluoromethyl; difluoromethyl; trifluoromethyl, C₁₋₄alkoxy and R¹ is of the formula A' or A'':

-(CH₂)_r



(A')

-CH₂CH(OH)(CH₂)_tCO₂H (A'')

wherein x is 0 or 1; r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

R² is hydrogen;

5

R³ is selected from C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyanoC₁₋₄alkyl, and C₁₋₄alkyl [substituted by 1 or 2 R⁸ groups (provided that when there are 2 R⁸ groups they are not substituents on the same carbon)];

10

{R⁸ is independently selected from hydroxy, heterocyclyl, C₁₋₄alkanoyl, C₁₋₄alkoxy, C₁₋₄alkanesulfinyl, C₁₋₄alkanesulfonyl, -COCOOR⁹, (R⁹)(R¹⁰)NCO-, -COCH₂OR¹¹, (R⁹)(R¹⁰)N-, -COOR⁹ and 2,2-dimethyl-1,3-dioxolan-4-yl;

15

[R⁹ and R¹⁰ are independently selected from hydrogen, hydroxy, C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C₁₋₄alkyl substituted by C₁₋₄alkoxy and wherein R⁹ and R¹⁰ can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

20

R¹¹ is selected from hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy and hydroxyc₁₋₄alkyl]}.
Another preferred class of compounds is of formula (1) wherein:

--- is a single bond;

X is N;

R⁴ and R⁵ are together -C(R⁷)=C(R⁶)-S-;

25

R⁶ is chloro;

R⁷ is hydrogen;

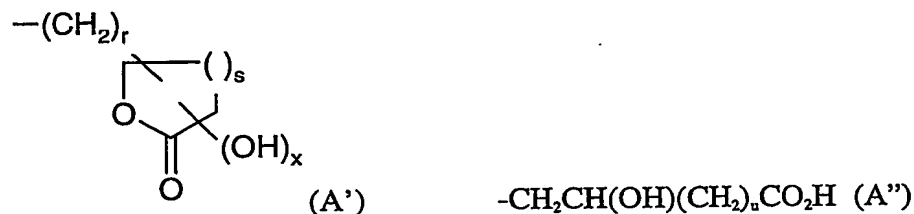
30

A is phenylene;

n is 1 or 2;

R^1 is independently selected from hydrogen, halo, nitro, hydroxy, C_{1-4} alkyl, C_{1-4} alkoxy and

R^1 is of the formula A' or A'' :



wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

R^2 is hydrogen;

R^3 is selected from C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano C_{1-4} alkyl, and C_{1-4} alkyl [substituted by 1 or 2 R^8 groups (provided that when there are 2 R^8 groups they are not substituents on the same carbon)];

15 { R^8 is independently selected from hydroxy, heterocyclyl, C_{1-4} alkanoyl, C_{1-4} alkoxy, C_{1-4} alkanesulfinyl, C_{1-4} alkylsulfonyl, $-COCOOR^9$, $(R^9)(R^{10})NCO-$, $-COCH_2OR^{11}$, $(R^9)(R^{10})N-$, $-COOR^9$ and 2,2-dimethyl-1,3-dioxolan-4-yl;

[R^9 and R^{10} are independently selected from hydrogen, hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C_{1-4} alkyl substituted by C_{1-4} alkoxy and wherein R^9 and R^{10} can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

25 R^{11} is selected from hydrogen, C_{1-4} alkyl, C_{1-4} alkoxy and hydroxy C_{1-4} alkyl].

Another preferred class of compound is of the formula (1) wherein:

---- is a single bond;

X is N;

R^4 and R^5 are together $-S-C(R^6)=C(R^7)-$;

R^6 is hydrogen or halo;

R^7 is hydrogen or halo;

5

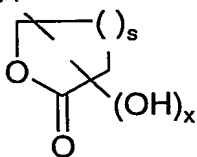
A is phenylene;

n is 1 or 2;

10 R^1 is independently selected from hydrogen, halo, nitro, hydroxy, C_{1-4} alkyl, C_{1-4} alkoxy and

R^1 is of the formula A' or A'':

$-(CH_2)_r$



(A')

$-CH_2CH(OH)(CH_2)_nCO_2H$ (A'')

wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not
15 a substituent on the ring carbon adjacent to the ring oxygen;

R^2 is hydrogen;

20 R^3 is selected from C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano C_{1-4} alkyl, and C_{1-4} alkyl [substituted by 1 or 2 R^8 groups (provided that when there are 2 R^8 groups they are not substituents on the same carbon)];

25 { R^8 is independently selected from hydroxy, heterocyclyl, C_{1-4} alkanoyl, C_{1-4} alkoxy, C_{1-4} alkanesulfinyl, C_{1-4} alkylsulfonyl, $-COCOOR^9$, $(R^9)(R^{10})NCO-$, $-COCH_2OR^{11}$, $(R^9)(R^{10})N-$, $-COOR^9$ and 2,2-dimethyl-1,3-dioxolan-4-yl;

[R^9 and R^{10} are independently selected from hydrogen, hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C_{1-4} alkyl substituted by C_{1-4} alkoxy and wherein R^9 and R^{10} can together with the nitrogen to which they are attached

form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

R^{11} is selected from hydrogen, C_{1-4} alkyl, C_{1-4} alkoxy and hydroxy C_{1-4} alkyl}.

5 A further preferred class of compound is of the formula (1) wherein;

--- is a single bond;

X is N;

R^4 and R^5 are together $-\text{C}(\text{R}^7)=\text{C}(\text{R}^6)-\text{S}-$;

10 R^6 is halo;

R^7 is hydrogen;

A is phenylene;

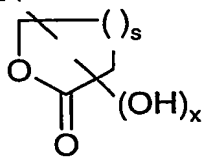
15

n is 1 or 2;

R^1 is independently selected from hydrogen, halo, nitro, hydroxy, C_{1-4} alkyl, C_{1-4} alkoxy and

20 R^1 is of the formula A' or A'':

$-(\text{CH}_2)_r$



$-\text{CH}_2\text{CH}(\text{OH})(\text{CH}_2)_u\text{CO}_2\text{H}$ (A'')

wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

25 R^2 is hydrogen;

R^3 is selected from C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon).

A further preferred class of compound is of the formula (1) wherein;

---- is a single bond;

X is N;

5 R^4 and R^5 are together $-C(R^7)=C(R^6)-S-$;

R^6 is halo;

R^7 is hydrogen;

10

A is phenylene;

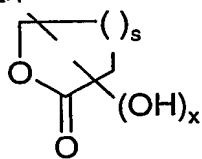
n is 1 or 2;

15

R^1 is independently selected from hydrogen, halo, nitro, hydroxy, C_{1-4} alkyl, C_{1-4} alkoxy and

R^1 is of the formula A' or A'':

$-(CH_2)_r$



(A')

$-CH_2CH(OH)(CH_2)_uCO_2H$ (A'')

wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not
20 a substituent on the ring carbon adjacent to the ring oxygen;

R^2 is hydrogen;

R^3 is selected from cyano C_{1-4} alkyl, and C_{1-4} alkyl substituted by R^8 ;

25

R^8 is selected from $(R^9)(R^{10})NCO-$, $(R^9)(R^{10})N-$, and $-COOR^9$;

R^9 and R^{10} are independently selected from hydrogen, hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C_{1-4} alkyl substituted by C_{1-4} alkoxy and wherein R^9 and R^{10} can together with the nitrogen to which they are attached

form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy.

Another class of compounds is of the formula (1) wherein

5 $\text{---}=\text{---}$ is a double bond;

X is N;

R^4 and R^5 are together $-\text{C}(\text{R}^7)=\text{C}(\text{R}^6)-\text{S}-$ or $-\text{S}-\text{C}(\text{R}^7)=\text{C}(\text{R}^6)-$;

R^6 is hydrogen or halo;

10

R^7 is hydrogen or hydrogen;

A is phenylene;

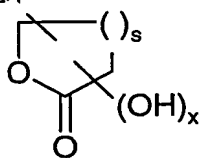
15

n is 1 or 2;

R^1 is independently selected from hydrogen, halo, nitro, hydroxy, C_{1-4} alkyl C_{1-4} alkoxy and

R^1 is of the formula A' or A'':

$-(\text{CH}_2)_r$



(A')

$-\text{CH}_2\text{CH}(\text{OH})(\text{CH}_2)_n\text{CO}_2\text{H}$ (A'')

20

wherein x is 0 or 1, r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

R^2 is carboxy;

25

R^3 is selected from hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano C_{1-4} alkyl, and C_{1-4} alkyl [substituted by 1 or 2 R^8 groups (provided that when there are 2 R^8 groups they are not substituents on the same carbon)];

{R⁸ is selected from hydroxy, C₁₋₄alkoxy, hydroxyC₁₋₄alkoxy, heterocyclyl, C₁₋₄alkanoyl, C₁₋₄alkoxy, C₁₋₄alkanesulfinyl, C₁₋₄alkylsulfonyl, -COCOOR⁹, (R⁹)(R¹⁰)NCO-, -COCH₂OR¹¹, (R⁹)(R¹⁰)N-, -COOR⁹ and 2,2-dimethyl-1,3-dioxolan-4-yl;

- 5 [R⁹ and R¹⁰ are independently selected from hydrogen, hydroxy, C₁₋₄alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C₁₋₄alkyl substituted by C₁₋₄alkoxy and wherein R⁹ and R¹⁰ can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

R¹¹ is selected from hydrogen, C₁₋₄alkyl, C₁₋₄alkoxy and hydroxyC₁₋₄alkyl]}.
 10

A further class of compound is of formula (1) wherein:

--- is a single bond;

- 15 X is N;

R⁴ and R⁵ are together -C(R⁷)=C(R⁶)-S- or -S-C(R⁷)=C(R⁶)-;

R⁶ is hydrogen or halo;

20

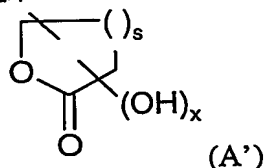
R⁷ is hydrogen or hydrogen;

A is pyridylene;

- 25 n is 1 or 2;

R¹ is independently selected from hydrogen, halo, nitro, hydroxy, C₁₋₄alkyl, C₁₋₄alkoxy and R¹ is of the formula A' or A'':

-(CH₂)_r



-CH₂CH(OH)(CH₂)_nCO₂H (A'')

wherein x is 0 or 1; r is 0, 1, 2 or 3 and s is 1 or 2; provided that the hydroxy group is not a substituent on the ring carbon adjacent to the ring oxygen;

R^2 is hydrogen;

5

R^3 is selected from hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon), cyano C_{1-4} alkyl, and C_{1-4} alkyl [substituted by 1 or 2 R^8 (provided that when there are 2 R^8 groups they are not substituents on the same carbon)];

10

{ R^8 is selected from hydroxy, C_{1-4} alkoxy, C_{1-4} alkoxy, hydroxy C_{1-4} alkoxy, heterocyclyl, C_{1-4} alkanoyl, C_{1-4} alkoxy, C_{1-4} alkanesulfinyl, C_{1-4} alkylsulfonyl, $-COCOOR^9$, $(R^9)(R^{10})NCO-$, $-COCH_2OR^{11}$, $(R^9)(R^{10})N-$, $-COOR^9$ and 2,2-dimethyl-1,3-dioxolan-4-yl};

15

[R^9 and R^{10} are independently selected from hydrogen, hydroxy, C_{1-4} alkyl (optionally substituted by 1 or 2 hydroxy groups provided that when there are 2 hydroxy groups they are not substituents on the same carbon) and C_{1-4} alkyl substituted by C_{1-4} alkoxy and wherein R^9 and R^{10} can together with the nitrogen to which they are attached form 4- to 6-membered ring where the ring is optionally substituted on carbon by 1 or 2 substituents selected from hydroxy or carboxy;

20

R^{11} is selected from hydrogen, C_{1-4} alkyl, C_{1-4} alkoxy and hydroxy C_{1-4} alkyl}.

25

In another aspect of the invention, preferred compounds of the invention are any one of:

methyl 2 [3-(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-

dihydroquinolin-1(2*H*)-yl]acetate;

2-[3-(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetic acid;

30

2-[3-(2-chloro-4*H*-thieno[3,2-*b*]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetic acid;

N-[1-(carbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-2-chloro-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;

- 2-chloro-*N*-[1-(dimethylcarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-[1-(methylcarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 5 2-chloro-*N*-[1-(*N*-hydroxycarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-{1-[(2-hydroxyethyl)carbamoylmethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-[1-(2-hydroxyethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 10 2-chloro-*N*-[1-(2,3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-2-chloro-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-{1-[(2,2-dimethyl-1,3-dioxolan-4(*S*)-yl)methyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl}-2-chloro-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 15 2-chloro-*N*-[1-(2(*S*),3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-[1-(2,2-Dimethyl-1,3-dioxolan-4(*R*)-ylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-[1-(2(*R*),3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 20 2-chloro-*N*-{1-[2-(4-hydroxypiperidin-1-yl)-2-oxoethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-{1-[*N*-(1,3-dihydroxyprop-2-yl)carbamoyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 25 2-chloro-*N*-{1-[*N*-(2-methoxyethyl)carbamoyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 2-chloro-*N*-(1-{2-[(3*a*,6*a*-*cis*)-2,2-dimethyltetrahydro-5*H*-[1,3]dioxolo[4,5-*c*]pyrrol-5-yl]-2-oxoethyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 30 2-chloro-*N*-(1-{2-[(*cis*)-3,4-dihydroxypyrrolidin-1-yl]-2-oxoethyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;

- methyl [3-(2,3-dichloro-4H-thieno[3,2-b]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2H)-yl]acetate;
- 2,3-Dichloro-N-[1-(2,3-dihydroxypropyl)-1,2,3,4-tetrahydro-2-oxo-3-quinoliny]-4H-thieno[3,2-b] pyrrole-5-carboxamide;
- 5 2,3-Dichloro-N-[1,2,3,4-tetrahydro-1-(2-hydroxyethyl)-2-oxo-3-quinoliny]-4H-thieno[3,2-b]pyrrole-5-carboxamide;
- 3-[[(2,3-Dichloro-4H-thieno[3,2-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-2-oxo-1(2H)-quinolineacetamide;
- 3-[[(2,3-Dichloro-4H-thieno[3,2-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-2-oxo-10 1(2H)-quinolineacetic acid;
- 2-Dichloro-N-[1-[2-[(3R,4S)-3,4-dihydroxy-1-pyrrolidinyl]-2-oxoethyl]-1,2,3,4-tetrahydro-2-oxo-3-quinoliny]- 6H-thieno[2,3-b]pyrrole-5-carboxamide;
- 3-[[(2,3-Dichloro-4H-thieno[3,2-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-2-oxo-, methyl ester 1(2H)-quinolineacetic acid;
- 15 3-[[(2-Chloro-6H-thieno[2,3-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-N-(2-methoxyethyl)-2- oxo-1(2H)-quinolineacetamide;
- 5-Chloro-N-[1-(2,3-dihydroxypropyl)-1,2,3,4-tetrahydro-2-oxo-3-quinoliny]-1H-thieno[2,3-d]imidazole-2- carboxamide
- 5,6-Dichloro-N-[1,2,3,4-tetrahydro-1-(2-hydroxyethyl)-2-oxo-3-quinoliny]-1H-thieno[2,3-d]imidazole-2-carboxamide
- 20 5-Chloro-N-[1,2,3,4-tetrahydro-1-(2-hydroxyethyl)-2-oxo-3-quinoliny]-1H-thieno[2,3-d]imidazole-2-carboxamide
- 5,6-Dichloro-N-[1-(2,3-dihydroxypropyl)-1,2,3,4-tetrahydro-2-oxo-3-quinoliny]-1H-thieno[2,3-d]imidazole-2-carboxamide

25

or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.

In another aspect of the invention, preferred compounds of the invention are any one of:

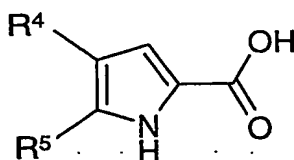
- 30 methyl 2 [3-(2-chloro-6H-thieno[2,3-b]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2H)-yl]acetate;
- 2-[3-(2-chloro-6H-thieno[2,3-b]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2H)-yl]acetic acid;

- 2-[3-(2-chloro-4*H*-thieno[3,2-*b*]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetic acid;
N-[1-(carbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-2-chloro-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 5 2-chloro-*N*-[1-(dimethylcarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]- 6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-[1-(methylcarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]- 6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-[1-(*N*-hydroxycarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]- 6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 10 2-chloro-*N*-{ 1-[(2-hydroxyethyl)carbamoylmethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-[1-(2-hydroxyethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]- 6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 15 2-chloro-*N*-[1-(2,3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-2-chloro-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-{ 1-[(2,2-dimethyl-1,3-dioxolan-4(*S*)-yl)methyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl}-2-chloro-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-[1-(2(*S*),3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 20 2-chloro-*N*-[1-(2,2-Dimethyl-1,3-dioxolan-4(*R*)-yl)methyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-[1-(2(*R*),3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 25 2-chloro-*N*-{ 1-[2-(4-hydroxypiperidin-1-yl)-2-oxoethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-{ 1-[*N*-(1,3-dihydroxyprop-2-yl)carbamoyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
2-chloro-*N*-{ 1-[*N*-(2-methoxyethyl)carbamoyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;
- 30 6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;

- 2-chloro-N-(1-{2-[(3a,6a-cis)-2,2-dimethyltetrahydro-5H-[1,3]dioxolo[4,5-c]pyrrol-5-yl]-2-oxoethyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6H-thieno[2,3-b]pyrrole-5-carboxamide;
- 2-chloro-N-(1-{2-[(cis)-3,4-dihydroxypyrrolidin-1-yl]-2-oxoethyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6H-thieno[2,3-b]pyrrole-5-carboxamide;
- 5 methyl [3-(2,3-dichloro-4H-thieno[3,2-b]pyrrol-5-yl)carbonylamino]-2-oxo-3,4-dihydroquinolin-1(2H)-yl]acetate;
- 2,3-Dichloro-N-[1-(2,3-dihydroxypropyl)-1,2,3,4-tetrahydro-2-oxo-3-quinolinyl]-4H-thieno[3,2-b]pyrrole-5-carboxamide;
- 10 2,3-Dichloro-N-[1,2,3,4-tetrahydro-1-(2-hydroxyethyl)-2-oxo-3-quinolinyl]-4H-thieno[3,2-b]pyrrole-5-carboxamide;
- 3-[[2,3-Dichloro-4H-thieno[3,2-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-2-oxo-1(2H)-quinolineacetamide;
- 3-[[2,3-Dichloro-4H-thieno[3,2-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-2-oxo-15 1(2H)-quinolineacetic acid;
- 2-Dichloro-N-[1-[2-[(3R,4S)-3,4-dihydroxy-1-pyrrolidinyl]-2-oxoethyl]-1,2,3,4-tetrahydro-2-oxo-3-quinolinyl]-6H-thieno[2,3-b]pyrrole-5-carboxamide;
- 3-[[2,3-Dichloro-4H-thieno[3,2-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-2-oxo-, methyl ester 1(2H)-quinolineacetic acid;
- 20 3-[[2-Chloro-6H-thieno[2,3-b]pyrrol-5-yl)carbonyl]amino]-3,4-dihydro-N-(2-methoxyethyl)-2-oxo-1(2H)-quinolineacetamide;
- or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.

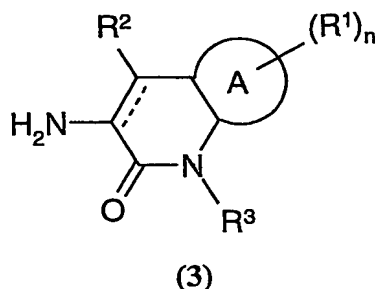
- Another aspect of the present invention provides a process for preparing a
- 25 compound of formula (1) or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof which process (wherein A, R¹, R², R³, R⁴, R⁵, n and --- are, unless otherwise specified, as defined in formula (1)) comprises of:

a) reacting an acid of the formula (2):



(2)

or an activated derivative thereof; with an amine of formula (3):



and thereafter if necessary:

- 5 i) converting a compound of the formula (1) into another compound of the formula (1);
- ii) removing any protecting groups;
- iii) forming a pharmaceutically acceptable salt or *in vivo* hydrolysable ester.

Specific reaction conditions for the above reaction are as follows.

- Process a)* Acids of formula (2) and amines of formula (3) may be coupled
- 10 together in the presence of a suitable coupling reagent. Standard peptide coupling reagents known in the art can be employed as suitable coupling reagents, or for example carbonyldiimidazole, 1-ethyl-3-(3-dimethylaminopropyl)carbodi-imide hydrochloride and dicyclohexyl-carbodiimide, optionally in the presence of a catalyst such as 1-hydroxybenzotriazole, dimethylaminopyridine or 4-pyrrolidinopyridine, optionally in the
 - 15 presence of a base for example triethylamine, di-isopropylethylamine, pyridine, or 2,6-di-alkyl-pyridines such as 2,6-lutidine or 2,6-di-*tert*-butylpyridine. Suitable solvents include dimethylacetamide, dichloromethane, benzene, tetrahydrofuran and dimethylformamide. The coupling reaction may conveniently be performed at a temperature in the range of -40 to 40°C.

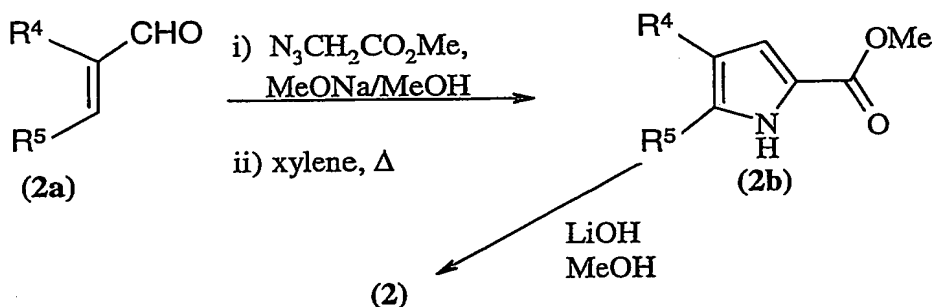
- 20 Suitable activated acid derivatives include acid halides, for example acid chlorides, and active esters, for example pentafluorophenyl esters. The reaction of these types of compounds with amines is well known in the art, for example they may be reacted in the presence of a base, such as those described above, and in a suitable solvent, such as those described above. The reaction may conveniently be performed at a
- 25 temperature in the range of -40 to 40°C.

Where R³ of formula (1) contains an ester group, the conversion of a compound of the formula (1) into another compound of the formula (1) may involve hydrolysis of the ester group. The reaction of this type is well known in the art.

Where R³ of formula (1) contains -COOH group, the conversion of a compound

of the formula (1) into another compound of the formula (1) may involve reduction of this group using reducing agents such as lithium borohydride and sodium borohydride. The conversion may also involve the coupling of this-COOH group with ammonia or a substituted amine in the presence of a base for example triethylamine, diisopropylethylamine, pyridine, or 2,6-di-*alkyl*-pyridines such as 2,6-lutidine or 2,6-di-*tert*-butylpyridine. Suitable solvents include dimethylacetamide, dichloromethane, benzene, tetrahydrofuran and dimethylformamide.

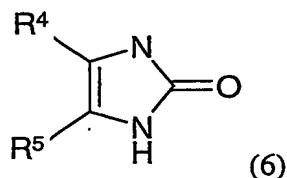
The acids of formula (2), wherein X is CH, may be prepared according to *Scheme 1*:



Scheme 1

Compounds of formula (2a) are commercially available or they are known compounds or they are prepared by processes known in the art.

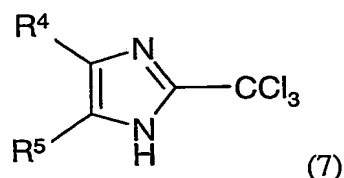
15 The acids of the formula (2), wherein X is N, can be prepared from a compound of the formula (6):



20 by firstly converting the oxo group to chlorine or bromine with a halogenating agent such as POCl₃ or POBr₃, in an inert organic solvent such as dichloromethane in a temperature range of ambient temperature to reflux (for example see *Nucleic Acid Chem.* 1991, 4, 24-6), then displacing the chlorine or bromine group with cyanide using a cyanide salt such as potassium cyanide, in an inert organic solvent such as toluene, benzene or xylene, 25 optionally in the presence of a catalyst such as 18-crown-6 (for example see *J.*

Heterocycl. Chem. 2000, 37(1), 119-126) and finally hydrolysing the cyano group to a carboxy group, with for example, an aqueous acid such as aqueous hydrogen chloride (for example see *Chem. Pharm. Bull.* 1986, 34(9), 3635-43).

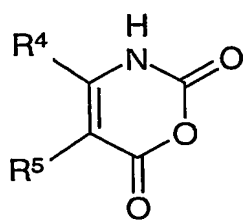
- 5 Alternatively, a compound of the formula (2) wherein X is N may be formed by reacting the compound of the formula (6) with $(\text{Cl}_3\text{CCO})_2\text{O}$ and $\text{Cl}_3\text{CCO}_2\text{H}$ in the presence of magnesium chloride using $\text{Cl}_3\text{CCO}_2\text{H}$ as solvent, to form a compound of the formula (7):



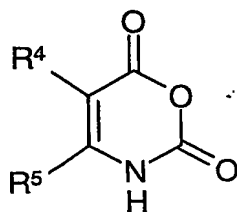
10

and then hydrolysing the compound of the formula (7), using, for example, aqueous sodium hydroxide, at a temperature range of ambient temperature to reflux (for example see *J Heterocycl. Chem.* 1980, 17(2), 381-2).

- 15 The compound of formula (6) may be prepared from a compound of formula (12) and (13) using conditions known for the Curtius rearrangement (*Tetrahedron* 1999, 55, 6167):

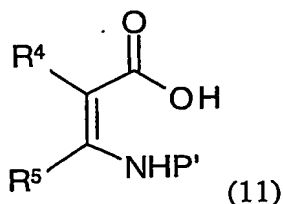
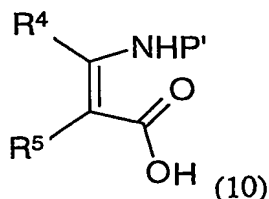


(12)



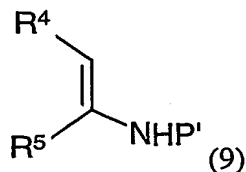
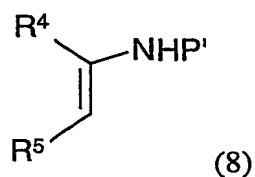
(13)

- 20 The compounds of the formula (10) and (11):



transform into compounds of the formula (12) and (13) respectively. This transformation either occurs spontaneously or may be induced with acid or base.

Compounds of the formula (10) and (11) may be prepared by introducing a carboxy group into a compound of the formula (8) or (9):

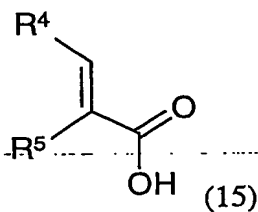
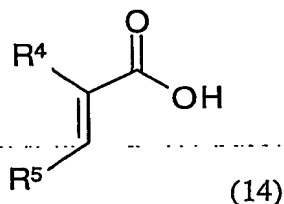


wherein P' is an amino protecting group such as butoxycarbonyl.

A carboxy group is introduced into the compound of the formula (8) or (9) by reacting an alkyl lithium reagent such as n-butyl lithium, in an inert organic solvent such as THF, at low temperature, for example in the range -10°C to -78°C and then forming the compound of the formula (10) or (11) as appropriate by either

- a) reacting the resulting compound with carbon dioxide; or
- b) by reacting with DMF in the temperature range of -10°C to ambient temperature to form the corresponding aldehyde and oxidizing the aldehyde to carboxy with standard reagents to give the compound of the formula (10) or (11).

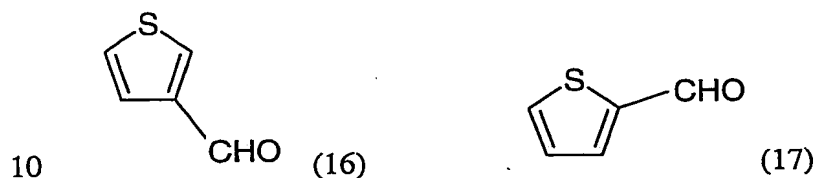
Compounds of the formula (8) and (9) may be prepared from a compound of the formula (14) and (15):



using conditions known for the Curtius reaction.

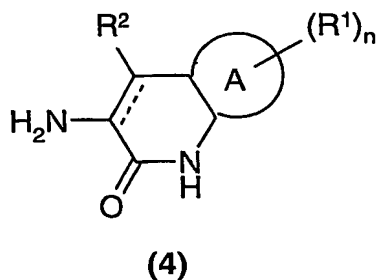
Compounds of the formula (14) and (15) may be prepared by oxidizing the corresponding aldehyde using standard oxidizing reagents such as potassium manganate or sodium periodate.

- 5 The aldehyde precursor of a compound of the formula (14) or (15) can be prepared using standard techniques known in the art. For example, many compounds of the formula (14) or (15) may be prepared by introducing the appropriate R^6 and R^7 into a compound of the formula (16) or (17) as appropriate:



- For example, when R^6 and R^7 are both chloro a compound of the formula (16) or (17) may be chlorinated with a chlorinating agent such as chlorine in the presence of aluminium chloride or iron (III) chloride, in an inert organic chlorinated solvent such as dichloromethane or 1,2-dichloroethane, followed by treatment with an aqueous base, such as, aqueous sodium hydroxide. The mono chlorinated compound can be formed in the same way.
- 15

Compounds of formula (3) may be prepared by reacting an amine of formula (4)



- 20 with R^3 -L where L is a suitable leaving group (for example chloro, bromo or iodo) in the presence of a base such as sodium hydride in a suitable solvent.

- It will be appreciated that certain of the various ring substituents in the compounds of the present invention may be introduced by standard aromatic substitution reactions or generated by conventional functional group modifications either prior to or immediately following the processes mentioned above, and as such are included in the
- 25

process aspect of the invention. Such reactions and modifications include, for example, introduction of a substituent by means of an aromatic substitution reaction, reduction of substituents, alkylation of substituents and oxidation of substituents. The reagents and reaction conditions for such procedures are well known in the chemical art. Particular
5 examples of aromatic substitution reactions include the introduction of a nitro group using concentrated nitric acid, the introduction of an acyl group using, for example, an acyl halide and Lewis acid (such as aluminium trichloride) under Friedel Crafts conditions; the introduction of an alkyl group using an alkyl halide and Lewis acid (such as aluminium trichloride) under Friedel Crafts conditions; and the introduction of a
10 halogen group. Particular examples of modifications include the reduction of a nitro group to an amino group by for example, catalytic hydrogenation with a nickel catalyst or treatment with iron in the presence of hydrochloric acid with heating; oxidation of alkylthio to alkylsulphinyl or alkylsulphonyl.

It will also be appreciated that in some of the reactions mentioned herein it may
15 be necessary/desirable to protect any sensitive groups in the compounds. The instances where protection is necessary or desirable and suitable methods for protection are known to those skilled in the art. Conventional protecting groups may be used in accordance with standard practice (for illustration see T.W. Green, Protective Groups in Organic Synthesis, John Wiley and Sons, 1991). Thus, if reactants include groups such as amino,
20 carboxy or hydroxy it may be desirable to protect the group in some of the reactions mentioned herein.

A suitable protecting group for an amino or alkylamino group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an alkoxycarbonyl group, for example a methoxycarbonyl, ethoxycarbonyl or *t*-butoxycarbonyl group, an
25 arylmethoxycarbonyl group, for example benzyloxycarbonyl, or an aroyl group, for example benzoyl. The deprotection conditions for the above protecting groups necessarily vary with the choice of protecting group. Thus, for example, an acyl group such as an alkanoyl or alkoxycarbonyl group or an aroyl group may be removed for example, by hydrolysis with a suitable base such as an alkali metal hydroxide, for example lithium or
30 sodium hydroxide. Alternatively an acyl group such as a *t*-butoxycarbonyl group may be removed, for example, by treatment with a suitable acid as hydrochloric, sulphuric or phosphoric acid or trifluoroacetic acid and an arylmethoxycarbonyl group such as a benzyloxycarbonyl group may be removed, for example, by hydrogenation over a catalyst

such as palladium-on-carbon, or by treatment with a Lewis acid for example boron tris(trifluoroacetate). A suitable alternative protecting group for a primary amino group is, for example, a phthaloyl group which may be removed by treatment with an alkylamine, for example dimethylaminopropylamine, or with hydrazine.

5 A suitable protecting group for a hydroxy group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an aroyl group, for example benzoyl, or an arylmethyl group, for example benzyl. The deprotection conditions for the above protecting groups will necessarily vary with the choice of protecting group. Thus, for example, an acyl group such as an alkanoyl or an aroyl group may be removed, for
10 example, by hydrolysis with a suitable base such as an alkali metal hydroxide, for example lithium or sodium hydroxide. Alternatively an arylmethyl group such as a benzyl group may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

 A suitable protecting group for a carboxy group is, for example, an esterifying
15 group, for example a methyl or an ethyl group which may be removed, for example, by hydrolysis with a base such as sodium hydroxide, or for example a *t*-butyl group which may be removed, for example, by treatment with an acid, for example an organic acid such as trifluoroacetic acid, or for example a benzyl group which may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

20 The protecting groups may be removed at any convenient stage in the synthesis using conventional techniques well known in the chemical art.

 Certain intermediates in the preparation of a compound of the formula (1) are novel and form another aspect of the invention.

 As stated hereinbefore the compounds defined in the present invention possesses
25 glycogen phosphorylase inhibitory activity. This property may be assessed, for example, using the procedure set out below.

Assay

 The activity of the compounds is determined by measuring the inhibitory effect of the compounds in the direction of glycogen synthesis, the conversion of glucose-1-
30 phosphate into glycogen with the release of inorganic phosphate, as described in EP 0 846 464 A2. The reactions were in 96well microplate format in a volume of 100µl. The change in optical density due to inorganic phosphate formation was measured at 620nm

in a Labsystems iEMS Reader MF by the general method of (Nordlie R.C and Arion W.J., Methods of Enzymology, 1966, 619-625). The reaction is in 50mM HEPES, 2.5mM MgCl₂, 2.25mM ethylene glycol-bis(b-aminoethyl ether) *N,N,N',N'*-tetraacetic acid, 100mM KCl, 2mM D-(+)-glucose pH7.2, containing 0.5mM dithiothreitol, the assay buffer solution, with 0.1mg type III glycogen, 0.15ug glycogen phosphorylase *a* (GP*a*) from rabbit muscle and 0.5mM glucose-1-phosphate. GP*a* is pre-incubated in the assay buffer solution with the type III glycogen at 2.5 mg ml⁻¹ for 30 minutes. 40μl of the enzyme solution is added to 25μl assay buffer solution and the reaction started with the addition of 25μl 2mM glucose-1-phosphate. Compounds to be tested are prepared in 10μl 10% DMSO in assay buffer solution, with final concentration of 1% DMSO in the assay. The non-inhibited activity of GP*a* is measured in the presence of 10μl 10% DMSO in assay buffer solution and maximum inhibition measured in the presence of 30μM CP320626 (Hoover et al (1998) J Med Chem 41, 2934-8; Martin et al (1998) PNAS 95, 1776-81). The reaction is stopped after 30min with the addition of 50μl acidic ammonium molybdate solution, 12ug ml⁻¹ in 3.48% H₂SO₄ with 1% sodium lauryl sulphate and 10ug ml⁻¹ ascorbic acid. After 30 minutes at room temperature the absorbency at 620nm is measured.

The assay is performed with a range of test concentrations of inhibitor to determine an IC₅₀, a concentration predicted to inhibit the enzyme reaction by 50%.

Activity is calculated as follows:-
% inhibition = $(1 - (\text{compound OD}_{620} - \text{fully inhibited OD}_{620}) / (\text{non-inhibited rate OD}_{620} - \text{fully inhibited OD}_{620})) * 100$.
OD₆₂₀ = optical density at 620nm.

Typical IC₅₀ values for compounds of the invention when tested in the above assay are in the range 100μM to 1nM. For example IC₅₀ of example 1 is 0.3μM.

According to a further aspect of the invention there is provided a pharmaceutical composition which comprises a compound of the formula (1), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, as defined hereinbefore in association with a pharmaceutically-acceptable diluent or carrier.

The composition may be in a form suitable for oral administration, for example as a tablet or capsule, for parenteral injection (including intravenous, subcutaneous, intramuscular, intravascular or infusion) as a sterile solution, suspension or emulsion, for

topical administration as an ointment or cream or for rectal administration as a suppository.

In general the above compositions may be prepared in a conventional manner using conventional excipients.

5 The compound of formula (1) will normally be administered to a warm-blooded animal at a unit dose within the range 5-5000 mg per square meter body area of the animal, i.e. approximately 0.1-100 mg/kg, and this normally provides a therapeutically-effective dose. A unit dose form such as a tablet or capsule will usually contain, for example 1-250 mg of active ingredient. Preferably a daily dose in the range
10 of 1-50 mg/kg is employed. However the daily dose will necessarily be varied depending upon the host treated, the particular route of administration, and the severity of the illness being treated. Accordingly the optimum dosage may be determined by the practitioner who is treating any particular patient.

15 According to a further aspect of the present invention there is provided a compound of the formula (1), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, as defined hereinbefore, for use in a method of treatment of a warm-blooded animal such as man by therapy.

20 According to an additional aspect of the invention there is provided a compound of the formula (1), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, as defined hereinbefore, for use as a medicament.

25 According to an additional aspect of the invention there is provided a compound of the formula (1), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, as defined hereinbefore, for use as a medicament in the treatment of type 2 diabetes, insulin resistance, syndrome X, hyperinsulinaemia, hyperglucagonaemia, cardiac ischaemia or obesity in a warm-blooded animal such as man.

30 According to this another aspect of the invention there is provided the use of a compound of the formula (1), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, as defined hereinbefore in the manufacture of a medicament for use in the treatment of type 2 diabetes, insulin resistance, syndrome X, hyperinsulinaemia, hyperglucagonaemia, cardiac ischaemia or obesity in a warm-blooded animal such as man.

According to this another aspect of the invention there is provided the use of a compound of the formula (1), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, as defined hereinbefore in the manufacture of a medicament for use in the treatment of type 2 diabetes in a warm-blooded animal such as man.

5 According to a further feature of this aspect of the invention there is provided a method of producing a glycogen phosphorylase inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (1).

10 According to this further feature of this aspect of the invention there is provided a method of treating type 2 diabetes, insulin resistance, syndrome X, hyperinsulinaemia, hyperglucagonaemia, cardiac ischaemia or obesity in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (1).

15 According to this further feature of this aspect of the invention there is provided a method of treating type 2 diabetes in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (1).

20 As stated above the size of the dose required for the therapeutic or prophylactic treatment of a particular cell-proliferation disease will necessarily be varied depending on the host treated, the route of administration and the severity of the illness being treated. A unit dose in the range, for example, 1-100 mg/kg, preferably 1-50 mg/kg is envisaged.

25 In addition to their use in therapeutic medicine, the compounds of formula (1) and their pharmaceutically acceptable salts are also useful as pharmacological tools in the development and standardisation of *in vitro* and *in vivo* test systems for the evaluation of the effects of inhibitors of cell cycle activity in laboratory animals such as cats, dogs, rabbits, monkeys, rats and mice, as part of the search for new therapeutic agents.

----- In the above other pharmaceutical composition, process, method, use and -----
medicament manufacture features, the alternative and preferred embodiments of the compounds of the invention described herein also apply.

Examples

The invention will now be illustrated by the following non-limiting examples in which, unless stated otherwise:

- 5 (i) temperatures are given in degrees Celsius ($^{\circ}\text{C}$); operations were carried out at room or ambient temperature, that is, at a temperature in the range of 18-25 $^{\circ}\text{C}$ and under an atmosphere of an inert gas such as argon;
- (ii) organic solutions were dried over anhydrous magnesium sulphate; evaporation of solvent was carried out using a rotary evaporator under reduced pressure (600-4000
10 Pascals; 4.5-30 mmHg) with a bath temperature of up to 60 $^{\circ}\text{C}$;
- (iii) chromatography means flash chromatography on silica gel; thin layer chromatography (TLC) was carried out on silica gel plates; where a Bond Elut column is referred to, this means a column containing 10 g or 20 g or 50 g of silica of 40 micron particle size, the silica being contained in a 60 ml disposable syringe and supported by a
15 porous disc, obtained from Varian, Harbor City, California, USA under the name "Mega Bond Elut SP"; "Mega Bond Elut" is a trademark; where a Biotage cartridge is referred to this means a cartridge containing KP-SILTM silica, 60 μ , particle size 32-63 μm , supplied by Biotage, a division of Dyax Corp., 1500 Avon Street Extended, Charlottesville, VA 22902, USA;
- 20 (iv) in general, the course of reactions was followed by TLC and reaction times are given for illustration only;
- (v) yields are given for illustration only and are not necessarily those which can be obtained by diligent process development; preparations were repeated if more material was required;
- 25 (vi) where given, NMR data is in the form of delta values for major diagnostic protons, given in parts per million (ppm) relative to tetramethylsilane (TMS) as an internal standard, determined at 300 MHz using perdeuterio dimethyl sulphoxide ($\text{DMSO}-d_6$) as solvent unless otherwise indicated, other solvents (where indicated in the text) include deuterated chloroform CDCl_3 ;
- 30 (vii) chemical symbols have their usual meanings; SI units and symbols are used;
- (viii) reduced pressures are given as absolute pressures in Pascals (Pa); elevated pressures are given as gauge pressures in bars;

(ix) solvent ratios are given in volume : volume (v/v) terms;

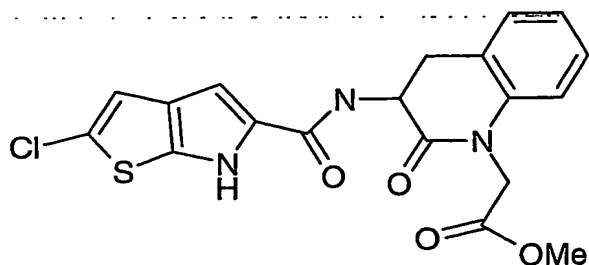
(x) mass spectra (MS) were run with an electron energy of 70 electron volts in the chemical ionisation (CI) mode using a direct exposure probe; where indicated ionisation was effected by electron impact (EI), fast atom bombardment (FAB) or electrospray (ESP); values for m/z are given; generally, only ions which indicate the parent mass are reported and unless otherwise stated the value quoted is $(M-H)^+$;

(xi) The following abbreviations are used:

	SM	starting material;
	EtOAc	ethyl acetate;
10	MeOH	methanol;
	EtOH	ethanol;
	DCM	dichloromethane;
	HOBt	1-hydroxybenzotriazole;
	DIPEA	di-isopropylethylamine;
15	EDCI	1-ethyl-3-(3-dimethylaminopropyl)carbodi-imide hydrochloride;
	Et ₂ O	diethyl ether;
	THF	tetrahydrofuran;
	DMF	<i>N, N</i> -dimethylformamide;
20	HATU	<i>O</i> -(7-Azabenzotriazol-1-yl)- <i>N, N, N', N'</i> - tetramethyluroniumhexafluorophosphate

Example 1

25 Methyl 2- 3-(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-yl)carbonylamino)-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetate

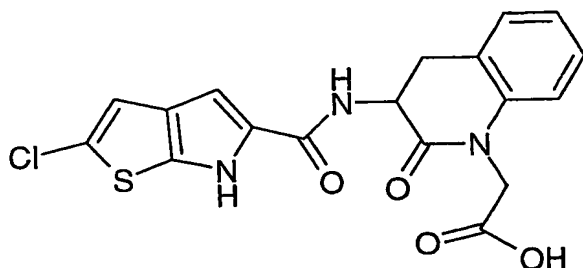


5-Carboxy-2-chloro-6*H*-thieno[2,3-*b*]pyrrole (**Method 9**; 5.07 g, 25.2 mmol), HOBt (3.40 g, 25.2 mmol), anhydrous DMF (100 mL) and finally EDCI (4.82 g, 25.2 mmol) were added to methyl 3-amino-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetate (**Method 1**; 5.89 g, 25.2 mmol) and the reaction was stirred for 18 h. The reaction was then diluted with water (200 mL) and stirred vigorously for 30 min. The resultant precipitate was filtered and washed with water (50 mL), EtOAc (2 x 20 mL) and Et₂O (2 x 10 mL). The collected solid was further dried under high vacuum for 6 h to furnish the title compound (8.00 g, 76%) as a pale yellow solid.

¹H NMR 3.15 (m, 2H), 3.64 (s, 3H), 4.74 (m, 3H), 7.18 (m, 6H), 8.58 (d, 1H), 11.91 (s, 1H); MS *m/z* MH⁺ 418, 420.

Example 2

2-[3-(2-Chloro-6*H*-thieno[2,3-*b*]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetic acid

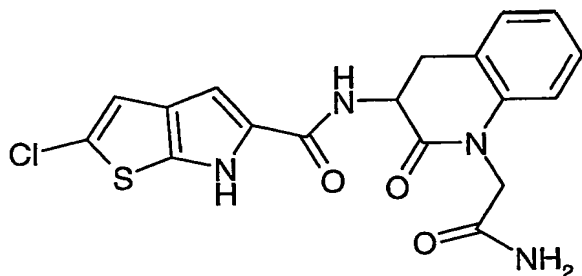


LiOH (1.41 g, 33.6 mmol) in H₂O (16.5 mL) was added to a stirring solution of (methyl 2-[3-(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetate) (**Example 1**; 7.00 g, 16.8 mmol) in THF (88 mL) and the reaction was stirred for 2 h. The reaction was quenched by addition of 1M aqueous HCl (200 mL) and EtOAc (400 mL) and the organic layer was dried (MgSO₄), filtered and evaporated. The resultant white foam was triturated with hot Et₂O (100 mL) cooled, filtered and dried to afford the title compound (6.00 g, 89%) as a white solid.

¹H NMR 3.14 (m, 2H), 4.52 (d, 1H), 4.75 (m, 2H), 7.03 (m, 3H), 7.18 (s, 1H), 7.27 (m, 2H), 7.57 (d, 1H), 11.90 (s, 1H), 12.89 (br. s, 1H); MS *m/z* MH⁺ 404, 406.

Example 3

2-Chloro-N-[1-(carbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide



5 Triethylamine (38 μ L, 0.27 mmol) then ethyl chloroformate (26.1 μ L, 0.27 mmol) were added to 2-[3-(2-chloro-6H-thieno[2,3-b]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2H)-yl]acetic acid (**Example 2**; 100 mg, 0.25 mmol) in anhydrous THF (2 mL) at 0 °C followed by stirring for 1 h. Concentrated aqueous NH_3 (1 mL) was
10 added and the reaction was stirred for a further 1 h. Water (20 mL) and EtOAc (40 mL) were added and the organic layer was separated, washed with 1M HCl (20 mL) and the organic layer was dried (MgSO_4), filtered and evaporated. The residue was purified by column chromatography (MeOH:DCM 1:19) to afford the title compound (56 mg, 56%) as a white solid.

15 ^1H NMR 3.15 (m, 2H), 4.23 (d, 1H), 4.67 (d, 1H), 4.82 (m, 1H), 6.88 (d, 1H), 7.05 (m, 2H), 7.14 (s, 2H), 7.24 (m, 2H), 7.54 (s, 1H), 8.51 (d, 1H), 11.91 (s, 1H); MS m/z MH^+ 403, 405.

Examples 4-7

20 The following examples were synthesised by an analogous method to **Example 3**:

Example 4

2-Chloro-N-[1-(N,N-dimethylcarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide

Example 5

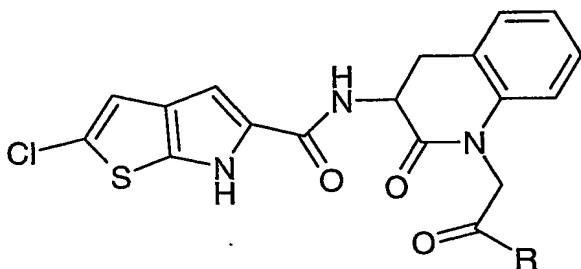
- 5 2-Chloro-N-[1-(N-methylcarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide




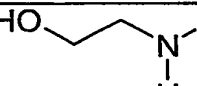
Example 6

2-Chloro-N-[1-(N-hydroxycarbamoylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide

- 10 **Example 7**

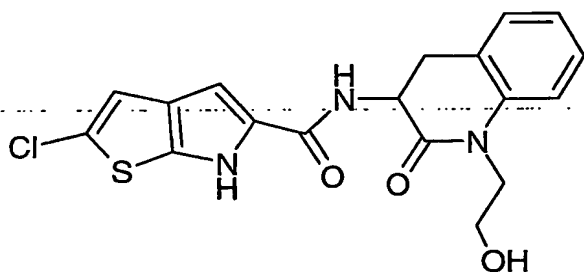
2-Chloro-N-{1-[N-(2-hydroxyethyl)carbamoylmethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6H-thieno[2,3-b]pyrrole-5-carboxamide



Example	NR ¹ R ²	¹ H NMR	m/z MH ⁺
4		2.84 (s, 3H), 3.03 (dd, 1H), 3.10 (s, 3H), 3.19 (app. t, 1H), 4.58 (d, 1H), 4.74 (m, 1H), 4.93 (d, 1H), 6.87 (d, 1H), 7.03 (t, 1H), 7.09 (s, 1H), 7.18 (s, 1H), 7.25 (m, 2H), 8.56 (d, 1H), 11.95 (s, 1H).	431, 433
5		2.60 (s, 3H), 3.05 (dd, 1H), 3.16 (app. t, 1H), 4.37 (d, 1H), 4.68 (d, 1H), 4.87 (m, 1H), 6.88 (d, 1H), 7.04 (t, 1H), 7.10 (s, 1H), 7.18 (s, 1H), 7.27 (m, 2H), 8.00 (m, 1H), 8.52 (d, 1H), 11.91 (s, 1H).	417, 419
6		3.02 (dd, 1H), 3.17 (app. t, 1H), 4.30 (d, 1H), 4.61 (d, 1H), 4.84 (m, 1H), 6.92 (d, 1H), 7.05 (t, 1H), 7.09 (s, 1H), 7.17 (s, 1H), 7.27 (m, 2H), 8.52 (d, 1H), 8.90 (s, 1H), 10.72 (s, 1H), 11.92 (s, 1H).	419, 421
7		3.04 (m, 1H), 3.10 (m, 3H), 3.39 (m, 2H), 4.32 (d, 1H), 4.73 (m, 3H), 6.89 (d, 1H), 7.03 (t, 1H), 7.10 (s, 1H), 7.17 (s, 1H), 7.27 (app. d, 2H), 8.09 (t, 1H), 8.53 (d, 1H), 11.92 (s, 1H).	447, 449

Example 8

- 5 2-Chloro-N-[1-(2-hydroxyethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6H-thieno[2,3-b]pyrrol-5-ylcarboxamide



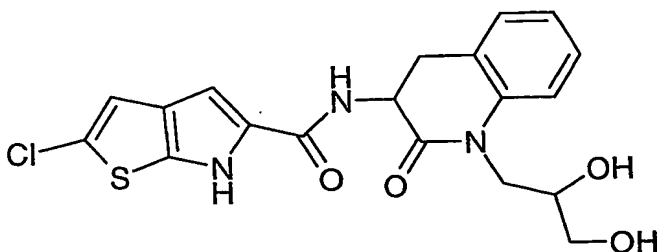
- Triethylamine (0.76 mL, 5.47 mmol) then ethyl chloroformate (0.52 mL, 5.47 mmol) were added to 2-[3-(2-chloro-6H-thieno[2,3-b]pyrrol-5-ylcarbonylamino)-2-oxo-1,2,3,4-dihydroquinolin-1(2H)-yl]acetic acid (Example 2; 2.0 g, 4.97 mmol) in anhydrous
- 10

THF (40 mL) at 0 °C followed by stirring for 1 h. LiBH₄ (2.0 M in THF, 3.1 mL, 6.21 mmol) was added slowly and the mixture stirred for a further 30 min. The reaction was carefully quenched with 1M HCl (200 mL) and EtOAc (400 mL) and the organic layer was further washed with sat. aqueous NaHCO₃ (100 mL), brine (100 mL), dried (MgSO₄), filtered and evaporated. The residue was triturated with refluxing Et₂O (30 mL) and after cooling the solid was filtered and dried to afford the title compound (1.70 g, 88%) a white solid.

¹H NMR 3.04 (m, 2H), 3.59 (m, 2H), 3.91 (m, 1H), 4.01 (m, 1H), 4.72 (m, 1H), 4.83 (m, 1H), 7.18 (m, 6H), 8.48 (d, 1H), 11.90 (s, 1H); MS m/z MH⁺ 390, 392.

Example 9

2-Chloro-N-[1-(2,3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide

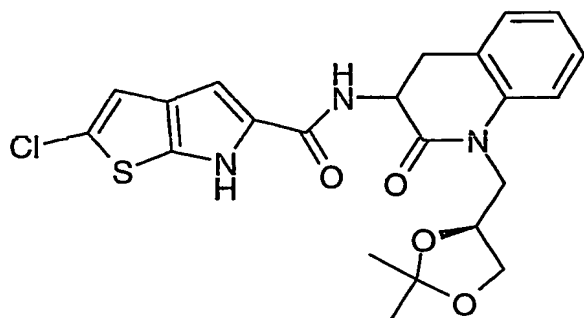


6M aqueous HCl (1.47 mL) was added to N-[1-(2,2-dimethyl-1,3-dioxolan-4-ylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl]-2-chloro-6H-thieno[2,3-b]pyrrole-5-carboxamide (Method 3; 340 mg, 7.45 mmol) in THF (14 mL) and the reaction was stirred for 4 h. The reaction was quenched by addition of triethylamine (1.5 mL) and the reaction was diluted with water (30 mL) and EtOAc (40 mL). The organic layer was separated, dried (MgSO₄), filtered and evaporated. The residue was triturated with hot Et₂O (10 mL) and after cooling was filtered and dried to afford the title compound (260 mg, 83%) as white solid.

¹H NMR 3.07 (m, 3H), 3.81 (m, 2H), 4.01 (m, 2H), 4.71 (m, 3H), 7.16 (m, 6H), 8.45 (app. d, 1H), 11.91 (s, 1H); MS m/z MH⁺ 420, 422.

Example 10

2-Chloro-N-[1-[(2,2-dimethyl-1,3-dioxolan-4(*S*)-yl)methyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide

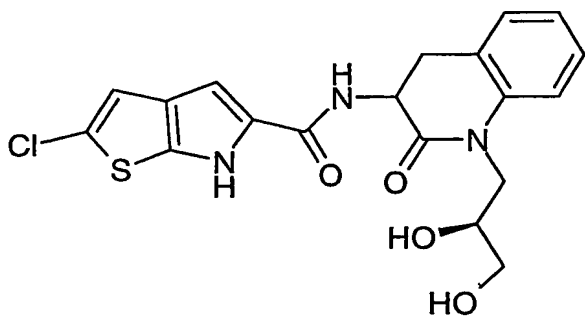


The title compound was prepared by exactly the same method as that described
 5 (Method 2) using [(4*R*)-2,2-dimethyl-1,3-dioxolan-4-yl]methyl methanesulfonate (*J. Med. Chem.*, 26, 1983 950-57), followed by the coupling procedure of Method 3.

¹H NMR 1.32 (s, 1.5H), 1.33 (s, 1.5H), 1.37 (s, 1.5H), 1.42 (s, 1.5H), 2.88 (m, 1H), 3.63 (m, 1H), 3.78 (app. t, 1H), 3.90 (dd, 0.5H), 4.04 (dd, 0.5H), 4.14 (m, 1H), 4.33 (m, 2H), 4.68 (m, 1H), 6.82 (m, 2H), 7.10 (m, 1H), 7.27 (m, 4H), 10.94 (br. s, 1H); MS *m/z* MNa⁺
 10 482, 484.

Example 11

2-Chloro-N-[1-(2(*S*),3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide

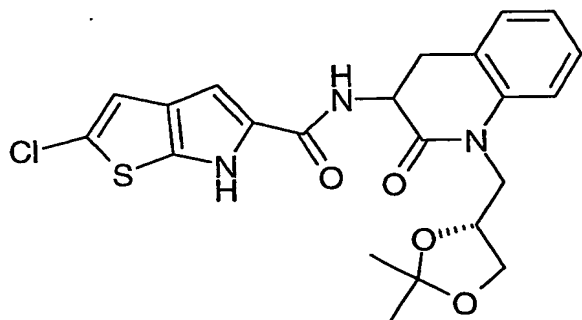


The title compound was prepared by acid hydrolysis by exactly the same method
 as that described (Example 9) starting with 2-chloro-N-[1-(2,2-dimethyl-1,3-dioxolan-4(*S*)-ylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(*R,S*)-yl]-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide (Example 10).

¹H NMR 3.07 (m, 3H), 3.81 (m, 2H), 4.01 (m, 2H), 4.71 (m, 3H), 7.16 (m, 6H), 8.45 (app. d, 1H), 11.91 (s, 1H); MS *m/z* MH⁺ 420, 422.

Example 12

2-Chloro-N-[1-(2,2-dimethyl-1,3-dioxolan-4(R)-ylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(R,S)-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide



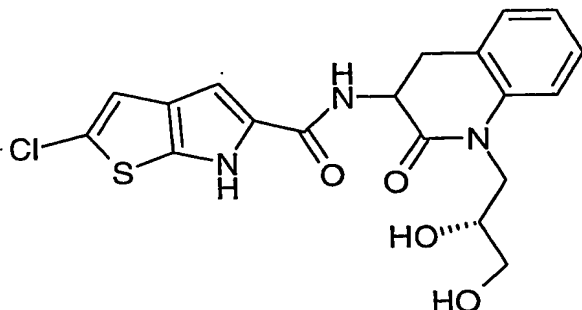
5

The title compound was prepared by exactly the same method as that described above (**Method 2**) using [(4*S*)-2,2-dimethyl-1,3-dioxolan-4-yl]methyl methanesulfonate (*J. Org. Chem.*, **64**, **1999** 6782-6790), followed by the coupling procedure of **Method 3**.
¹H NMR 1.32 (s, 1.5H), 1.33 (s, 1.5H), 1.37 (s, 1.5H), 1.42 (s, 1.5H), 2.88 (m, 1H), 3.63 (m, 1H), 3.78 (app. t, 1H), 3.90 (dd, 0.5H), 4.04 (dd, 0.5H), 4.14 (m, 1H), 4.33 (m, 2H), 4.68 (m, 1H), 6.82 (m, 2H), 7.10 (m, 1H), 7.27 (m, 4H), 10.94 (br. s, 1H); MS *m/z* MNa⁺ 482, 484.

10

Example 13

2-Chloro-N-[1-(2(R),3-dihydroxypropyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(R,S)-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide



15

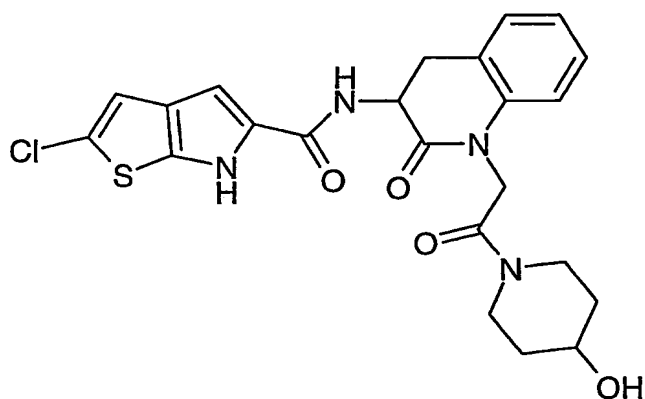
20

The title compound was prepared by acid hydrolysis by exactly the same method as that described above (**Example 9**) starting with 2-chloro-N-[1-(2,2-dimethyl-1,3-dioxolan-4(R)-ylmethyl)-2-oxo-1,2,3,4-tetrahydroquinolin-3(R,S)-yl]-6H-thieno[2,3-b]pyrrole-5-carboxamide (**Example 12**).

¹H NMR 3.07 (m, 3H), 3.81 (m, 2H), 4.01 (m, 2H), 4.71 (m, 3H), 7.16 (m, 6H), 8.45 (app. d, 1H), 11.91 (s, 1H); MS m/z MH⁺ 420, 422.

5 Example 14

2-Chloro-N-{1-[2-(4-hydroxypiperidin-1-yl)-2-oxoethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-6H-thieno[2,3-b]pyrrole-5-carboxamide



10

4-Dimethylaminopyridine (5 mg, 0.038mmol) and 4-hydroxypiperidine (42 mg, 0.41 mmol) were added to a suspension of 2-[3-(2-Chloro-6H-thieno[2,3-b]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2H)-yl]acetic acid (**Example 2**; 150 mg, 0.38 mmol) and EDCI (79 mg, 0.41 mmol) in THF (0.5 mL) under an inert atmosphere.

15 DMF (0.5 mL) was added and the mixture stirred at ambient temperature for 18 h. After pouring into water (10 mL) the resultant solid was filtered off and washed with 1M HCl aq. and water. Chromatography on silica gel (eluent gradient of CH₂Cl₂ to MeOH:CH₂Cl₂ (1:9)) afforded the title compound (109 mg, 59%) as an off white solid.

20 ¹H NMR (400MHz) 1.20-1.52 (m, 2H), 1.65-1.90 (m, 2H), 3.05 (m, 2H), 3.27(m, 2H), 3.80 (m, 3H), 4.64 (dd, 1H), 4.75 (m, 2H), 4.96 (dd, 1H), 6.89 (d, 1H), 7.04 (t, 1H), 7.11 (s, 1H), 7.19 (s, 1H), 7.28 (t, 1H), 8.54 (d, 1H), 11.93 (s, 1H); MS m/z MH⁺ 487, 489

Examples 15-17

The following examples were synthesised by an analogous method to **Example**

25 **14:**

Example 15

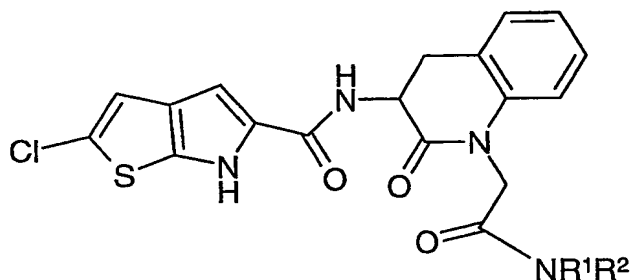
2-Chloro-*N*-(1-[*N*-(1,3-dihydroxyprop-2-yl)carbamoylmethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide

Example 16

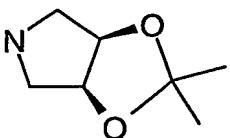
- 5 2-chloro-*N*-(1-[*N*-(2-Methoxyethyl)carbamoylmethyl]-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide

Example 17

- 10 2-chloro-*N*-(1-{2-[(3*a*,6*a*-*cis*)-2,2-dimethyltetrahydro-5*H*-[1,3]dioxolo[4,5-*c*]pyrrol-5-yl]-2-oxoethyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide

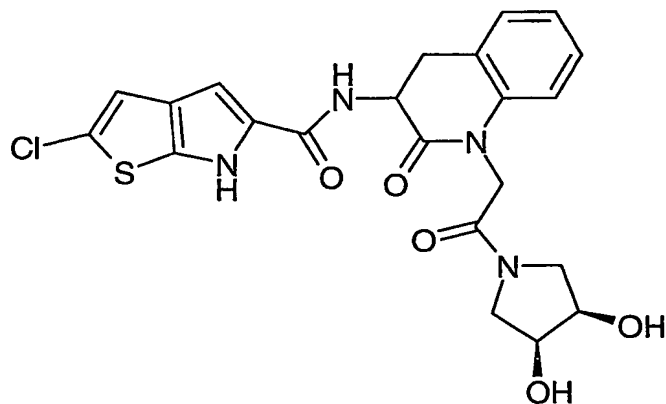


Example	NR¹R²	¹ H NMR (400MHz)	m/z	SM (amine)
15		3.05 (dd, 1H), 3.18 (dd, 1H), 3.42 (m, 4H), 3.73 (m, 1H), 4.40 (d, 1H), 4.63 (m, 2H), 4.77 (m, 2H), 6.93 (d, 1H), 7.05 (t, 1H), 7.11 (s, 1H), 7.19 (s, 1H), 7.28 (t, 1H), 7.88 (d, 1H), 8.55 (d, 1H), 11.94 (s, 1H).	477, 479	commercial
16		3.05 (dd, 1H), 3.25 (m, 9H), 4.33 (d, 1H), 4.73 (d, 1H), 4.85 (m, 1H), 6.90 (d, 1H), 7.06 (t, 1H), 7.11 (s, 1H), 7.19 (s, 1H), 7.29 (t, 1H), 8.20 (t, 1H), 8.54	461, 463	commercial

		(d, 1H), 11.94 (s, 1H).		
17		1.28, 1.29, 1.39, 1.42 (4 x s, 6H), 3.05 (dd, 1H), 3.19 (m, 2H), 3.61 (quintet, 1H), 3.74 (dd, 1H), 3.88 (dd, 1H), 4.59 (dd, 1H), 4.84 (m, 4H), 6.89 (t, 1H), 7.05 (t, 1H), 7.11 (s, 1H), 7.19 (s, 1H), 7.24 (t, 1H), 7.30 (d, 1H), 8.56 (d, 1H), 11.94 (s, 1H).	529, 531	Org.Lett., 3, 2001 465-468

Example 18

5 2-Chloro-N-(1-{2-[(*cis*)-3,4-Dihydroxypyrrolidin-1-yl]-2-oxoethyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6H-thieno[2,3-*b*]pyrrole-5-carboxamide



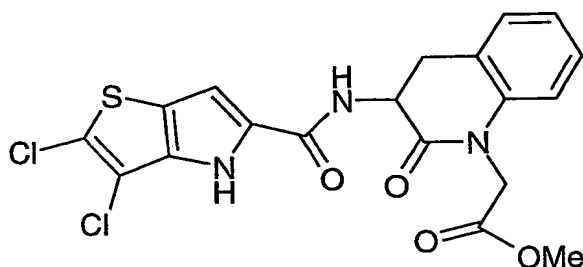
1M HCl aq. (0.46 mL, 0.46 mmol) was added to 2-chloro-N-(1-{2-[(3a,6a-*cis*)-2,2-dimethyltetrahydro-5H-[1,3]dioxolo[4,5-*c*]pyrrol-5-yl]-2-oxoethyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl)-6H-thieno[2,3-*b*]pyrrole-5-carboxamide (**Example 17**; 200 mg, 0.38 mmol) in EtOH (6 mL) and heated to 70 °C for 3 hrs. 2M HCl aq. (1 mL) was added and the mixture heated to 70°C for 18 h. After cooling all volatiles were removed under reduced pressure. Chromatography on silica gel (eluent gradient of CH₂Cl₂ to THF) and washing the resultant solid with Et₂O then hexane afforded the title compound (170 mg, 92%) as an off white solid.

^1H NMR (400MHz) 3.05 (dd, 1H); 3.21 (m, 2H), 3.42 (m, 2H), 3.77 (dd, 1H), 4.03 (m, 1H), 4.14 (m, 1H), 4.51 (dd, 1H), 4.77 (m, 2H), 4.92 (d, 1H), 5.02 (d, 1H), 6.93 (d, 1H), 7.05 (t, 1H), 7.11 (s, 1H), 7.19 (s, 1H), 7.28 (m, 2H), 8.55 (d, 1H), 11.94 (s, 1H); MS m/z MH^+ 489, 491.

5

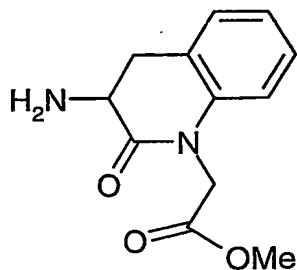
Example 19

Methyl [3-(2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrol-5-ylcarbonylamino)-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl]acetate



10 5-Carboxy-2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrole (**Method 8**; 595 mg, 2.52 mmol), HOBt (340 mg, 2.52 mmol), DCM (100 mL) and finally EDCI (483 mg, 2.52 mmol) were added to methyl (3-amino-2-oxo-3,4-dihydroquinolin-1(2*H*)-yl)acetate (**Method 1**, 590 mg, 2.52 mmol) and the reaction was stirred for 18 h. The reaction was then diluted with water (50 mL) and stirred vigorously for 30 min. The resultant
15 precipitate was filtered and washed with Et₂O (2 x 20 mL). After filtration the resultant solid was then triturated with refluxing Et₂O (25 mL) and after cooling the title compound (528 mg, 46%) was collected again by filtration as a white solid.
 ^1H NMR 3.10 (dd, 1H), 3.21 (app. t, 1H), 3.69 (s, 3H), 4.67 (d, 1H), 4.81 (m, 2H), 7.07 (m, 2H), 7.23 (s, 1H), 7.31 (m, 2H), 8.69 (d, 1H), 12.51 (s, 1H); MS m/z MH^+ 452, 454.

20

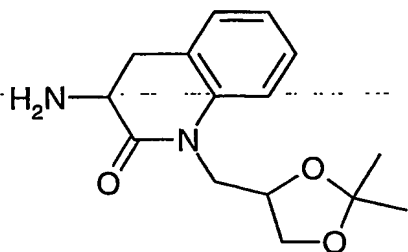
MethodsMethod 1Methyl (3-amino-2-oxo-3,4-dihydroquinolin-1(2H)-yl)acetate

5

Sodium hydride (60% in oil, 2.52 g, 63.0 mmol) was added to 3-amino-3,4-dihydroquinolin-2(1H)-one hydrochloride (*J. Med. Chem.*, 28, 1985, 1511-16; 5.0 g, 25.2 mmol), in anhydrous DMF (100 mL) at 0 °C over a period of 5 min keeping the internal temperature at <10 °C. The reaction was stirred for a further 30 min before addition of methyl bromoacetate (2.85 mL, 30.2 mmol), then stirred for a further 60 min. The reaction was quenched by addition of 1M aqueous HCl (5 mL) and the volatiles were removed by evaporation. The residue was dissolved in DCM (250 mL) and washed with sat. aqueous NaHCO₃ (100 mL) and the organic layer was dried (MgSO₄), filtered and evaporated to yield the title compound (5.89 g, 25.2 mmol) as yellow paste which was used without further purification.

10
15

¹H NMR 2.21 (br. s, 2H), 2.78 (d, 1H), 2.97 (dd, 1H), 3.47 (dd, 1H), 3.67 (s, 3H), 4.55 (d, 1H), 4.78 (d, 1H), 6.96 (m, 2H), 7.23 (m, 2H); MS m/z MH⁺ 235.

Method 220 3-Amino-1-(2,2-dimethyl-1,3-dioxolan-4-ylmethyl)-3,4-dihydroquinolin-2(1H)-one

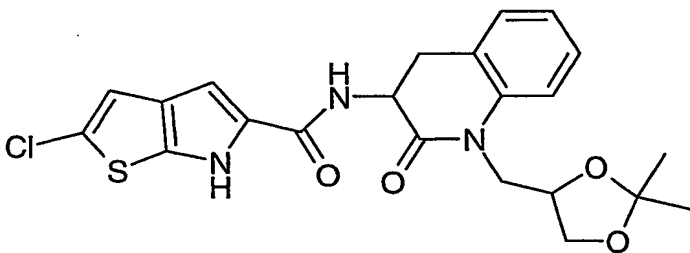
Sodium hydride (60% in oil, 191 mg, 4.70 mmol) was added to 3-amino-3,4-dihydroquinolin-2(1H)-one hydrochloride (*J. Med. Chem.*, 28, 1985; 1511-16, 450 mg, 2.27 mmol), in anhydrous DMF (6 mL) at 0 °C over a period of 5 min keeping the

internal temperature at $<10^{\circ}\text{C}$. The reaction was stirred for a further 30 min before addition of [2,2-dimethyl-1,3-dioxolan-4-yl]methyl methanesulfonate (*J. Med. Chem.* 35, 1992, 1650-62; 528 mg, 2.50 mmol) and the reaction has then heated to 80°C for a period of 5 h. The reaction was then cooled and evaporated before addition of sat. aqueous NaHCO_3 (20 mL) and EtOAc (50 mL). The organic layer was then dried (MgSO_4), filtered and evaporated and the residue was purified by column chromatography (MeOH:DCM 1:19) to afford the title compound (330 mg, 53%) as colourless oil.

^1H NMR 1.33 (s, 3H), 1.40 (s, 1.5H), 1.45 (s, 1.5H), 1.96 (br. s, 2H), 2.89 (m, 1H), 3.07 (m, 1H), 3.60 (m, 1H), 3.82 (m 1.5H), 4.08 (m, 1.5H), 4.33 (m, 2H), 7.04 (m, 1H), 7.23 (m, 3H); MS m/z MH^+ 277.

Method 3

N-{1-(2,2-Dimethyl-1,3-dioxolan-4-yl)methyl}-2-oxo-1,2,3,4-tetrahydroquinolin-3-yl}-2-chloro-6H-thieno[2,3-b]pyrrol-5-ylcarboxamide



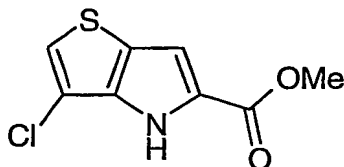
5-Carboxy-2-chloro-6H-thieno[2,3-b]pyrrole (**Method 9**; 243 mg, 1.20 mmol), HOBt (178 mg, 1.32 mmol), anhydrous DMF (10 mL) and finally EDCI (252 mg, 1.32 mmol) were added to 3-amino-1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-3,4-dihydroquinolin-2(1H)-one (**Method 2**, 330 mg, 1.20 mmol) and the reaction was stirred for 18 h. The reaction was evaporated and the residue was dissolved in EtOAc (100 mL) and washed with 1M aqueous HCl (50 mL) and the organic layer was further washed with sat. aqueous NaHCO_3 (30 mL) and brine (30 mL). The organic layer was then separated, dried (MgSO_4), filtered and evaporated. The residue was purified by column chromatography (EtOAc:hexanes 1:2) to afford the title compound (382 mg, 69%) as a white solid.

^1H NMR 1.32 (s, 1.5H), 1.33 (s, 1.5H), 1.37 (s, 1.5H), 1.42 (s, 1.5H), 2.88 (m, 1H), 3.63 (m, 1H), 3.78 (app. t, 1H), 3.90 (dd, 0.5H), 4.04 (dd, 0.5H), 4.14 (m, 1H), 4.33 (m, 2H),

4.68 (m, 1H), 6.82 (m, 2H), 7.10 (m, 1H), 7.27 (m, 4H), 10.94 (br. s, 1H); MS m/z MNa⁺ 482, 484.

Method 4

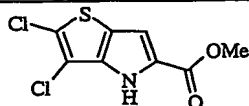
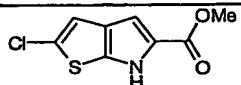
5 3-Chloro-5-methoxycarbonyl-4H-thieno[3,2-b]pyrrole



Methanolic sodium methoxide solution (28%) (5 ml, 25.9 mmol) was diluted with MeOH (5 ml) and was cooled to -25°C under nitrogen. A solution of 4-chloro-2-thienylcarboxaldehyde (J Heterocyclic Chem, 1976, 13, 393; 1.1 g, 7.5 mmol) and methyl azidoacetate (3.0 g, 26.1 mmol) in MeOH (20 ml) was added dropwise, maintaining the temperature at -25°C. On completion of addition the solution was allowed to warm to 5°C over a period of approximately 16 hours. The solution was added to saturated aqueous ammonium chloride (250 ml) and the mixture was extracted using DCM. The combined organic layers were concentrated at 0°C. The residue was taken up in xylene (30 ml) and this solution was added dropwise to xylene (120 ml) under reflux. The solution was heated under reflux for 30 minutes before being cooled and concentrated. The title compound was purified by a mixture of crystallisation (EtOAc/isohexane) and chromatography on a Bond Elut column eluting with a graduated solvent of 5-50% EtOAc in isohexane (640 mg, 40%). NMR (CDCl₃) 9.1 (1H, br), 7.1 (2H, s), 3.9 (3H, s); m/z 214.3.

Methods 5 and 6

The following compounds were made by the process of Method 4 using the appropriate starting materials

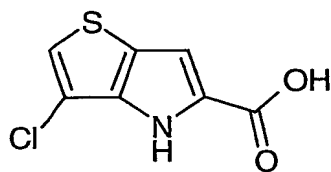
Meth.	Compound	NMR (CDCl ₃)	M/z
5 ¹		9.2 (1H, br), 7.0 (1H, s), 3.9 (3H, s)	248.2
6 ²		9.4-9.2 (1H, br), 7.0 (1H, s), 6.9 (1H, s), 3.9 (3H, s)	214

1 Aldehyde: DE 2814798

2 Aldehyde: Aldehyde ref. Gronowitz *et al.* Tetrahedron Vol.32 1976 p.1403

5 Method 7

5-Carboxy-3-chloro-4H-thieno[3,2-b]pyrrole

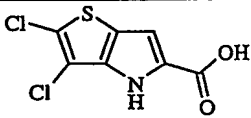
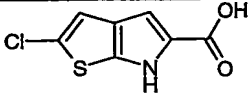


3-Chloro-5-methoxycarbonyl-4H-thieno[3,2-b]pyrrole (Method 4; 0.61 g, 2.83 mmol) was taken up in MeOH (10 ml) and was heated under reflux. Aqueous lithium hydroxide (2.0 M, 3.0 ml, 6.0 mmol) was added portionwise over 45 minutes. The mixture was heated under reflux for 30 minutes before being cooled and concentrated. Water (20 ml) was added and the solution was neutralised using aqueous hydrochloric acid (2.0 M, 3.0 ml). The solution was extracted using EtOAc, and the combined organic layers were concentrated to afford the title compound as a yellow solid (0.57 g, 100%).

NMR: 12.4 (1H, br), 7.4 (1H, s), 7.0 (1H, s); m/z 200.3.

Methods 7 and 8

The following compounds were made by the process of **Method 7** using the appropriate starting materials.

Method	Compound	NMR	M/z	SM
8		7.0 (1H, s)	234.2	Method 5
9		12.6-12.7 (1H, b), 12.0-12.1 (1H, b), 7.15 (1H, s), 6.9 (1H, s)	183	Method 6

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